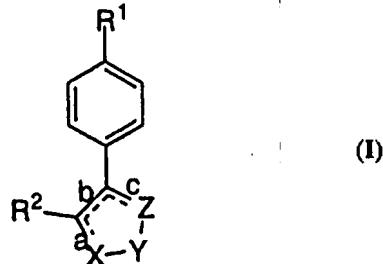




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : C07D 277/02, 275/02, 307/02, 333/04 A61K 31/33, C07C 305/18, 307/02		A3	(11) International Publication Number: WO 95/00501 (43) International Publication Date: 5 January 1995 (05.01.95)
(21) International Application Number: PCT/CA94/00318		[CA/CA]: 13199 Edison Crescent, Pierrefonds, Quebec H8Z 1Y5 (CA). LEGER, Serge [CA/CA]; 51 Lamarche, Dollard des Ormeaux, Quebec H9B 3E5 (CA). THERIEN, Michel [CA/CA]; 944 21st Avenue, Laval, Quebec H7R 2R2 (CA).	
(22) International Filing Date: 9 June 1994 (09.06.94)		(74) Agent: MURPHY, Kevin, P.; Swabey, Ogilvy, Renault, Suite 800, 1001 de Maisonneuve Boulevard West, Montreal, Quebec H3A 3C8 (CA).	
(30) Priority Data: 082,196 24 June 1993 (24.06.93) US 179,467 10 January 1994 (10.01.94) US		(81) Designated States: AU, BB, BG, BR, BY, CA, CN, CZ, FI, GE, HU, JP, KE, KG, KR, KZ, LK, LV, MD, MG, MN, MW, NO, NZ, PL, RO, RU, SD, SI, SK, TJ, TT, UA, US, UZ, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).	
(60) Parent Application or Grant (63) Related by Continuation US Filed on 179,467 (CIP) 10 January 1994 (10.01.94)		Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.	
(71) Applicant (for all designated States except US): MERCK FROSST CANADA INC. [CA/CA]; 16711 Trans-Canada Highway, Kirkland, Quebec H9H 3L1 (CA).		(88) Date of publication of the international search report: 13 April 1995 (13.04.95)	
(54) Title: PHENYL HETEROCYCLES AS CYCLOOXYGENASE-2 INHIBITORS			
(57) Abstract			
<p>The invention encompasses the novel compound of formula (I) useful in the treatment of cyclooxygenase-2 mediated diseases. The invention also encompasses certain pharmaceutical compositions for treatment of cyclooxygenase-2 mediated diseases comprising compounds of formula (I).</p>			



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A. CLASSIFICATION OF SUBJECT MATTER
IPC 5 C07D277/02 C07D275/02 C07D307/02 C07D333/04 A61K31/33
C07C305/18 C07C307/02

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 5 C07D A61K C07C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO,A,91 19708 (FUJISAWA PHARMACEUTICAL CO.) 26 December 1991 see examples ---	1-33
E X	WO,A,94 15932 (SEARLE & CO.) 21 July 1994 * see example 13 *	1-33 1-22, 24-33
X	* see scheme IV, compound 15, p. 32 ; see example 13, step 4 *	23
Y	EP,A,0 087 629 (DU PONT DE NEMOURS) 7 September 1983 see pages 13-14 see page 8; table I ---	1-33
		-/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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- *&* document member of the same patent family

5

Date of the actual completion of the international search

Date of mailing of the international search report

25 January 1995

07.03.95

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentiaan 2
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Fax (+ 31-70) 340-3016

Authorized officer

Lauro, P

C(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	CHEMICAL ABSTRACTS, vol. 113, no. 21, 19 November 1990, Columbus, Ohio, US; abstract no. 184393m, page 33 ; & J. PHARMACOL. EXP. THER., vol.254, no.1, 1990 pages 180 - 187 GANS,K; GALBRAITH,W; ROMAN R. ET AL. 'Anti-inflammatory and safety profile of DuP 697, a novel orally effective prostaglandin synthesis inhibitor' * see abstract * -----	1-33
Y	EP,A,0 388 909 (FUJISAWA PHARMACEUTICAL CO.) 26 September 1990 see claim 1 see page 9 - page 11 -----	1-33

INTERNATIONAL SEARCH REPORT

Internal application No.

PCT/CA94/00318

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:

2. Claims Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

Please see attached sheet ./.

3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.

2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest

No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/

iii) Incomplete search

In the light of the variety of (hetero)cyclic rings and the nature of their substituents, a meaningful search of the subject-matter of the claims 1-18, 27-33 could not be carried out (Article 17 (2)(a)(ii) PCT). The search has been limited to the examples, i.e. to the exemplified heterocyclic rings substituted with an additional phenyl ring or cyclohexyl adjycnt to the sulfonyl-substituted phenyl.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/LA 94/00318

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO-A-9119708	26-12-91	AU-A-	7973191	07-01-92
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		JP-A-	3027370	05-02-91
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		US-A-	5217971	08-06-93
		US-A-	5229386	20-07-93
		CA-A-	2012716	22-09-90



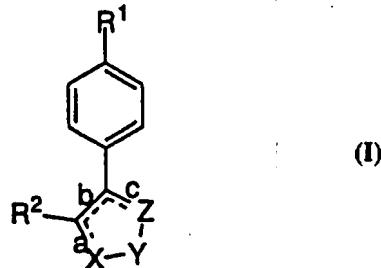
INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁵ : C07D 277/02, 275/02, 307/02, 333/04, A61K 31/00		A2	(11) International Publication Number: WO 95/00501 (43) International Publication Date: 5 January 1995 (05.01.95)
(21) International Application Number: PCT/CA94/00318			[CA/CA]; 13199 Edison Crescent, Pierrefonds, Quebec H8Z 1Y5 (CA). LEGER, Serge [CA/CA]; 51 Lamarche, Dollard des Ormeaux, Quebec H9B 3E5 (CA). THERIEN, Michel [CA/CA]; 944 21st Avenue, Laval, Quebec H7R 2R2 (CA).
(22) International Filing Date: 9 June 1994 (09.06.94)			
(30) Priority Data: 082,196 24 June 1993 (24.06.93) 179,467 10 January 1994 (10.01.94)		US US	(74) Agent: MURPHY, Kevin, P.; Swabey, Ogilvy, Renault, Suite 800, 1001 de Maisonneuve Boulevard West, Montreal, Quebec H3A 3C8 (CA).
(60) Parent Application or Grant (63) Related by Continuation US 179,467 (CIP) Filed on 10 January 1994 (10.01.94)			(81) Designated States: AU, BB, BG, BR, BY, CA, CN, CZ, FI, GE, HU, JP, KE, KG, KR, KZ, LK, LV, MD, MG, MN, MW, NO, NZ, PL, RO, RU, SD, SI, SK, TJ, TT, UA, US, UZ, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).
(71) Applicant (for all designated States except US): MERCK FROSST CANADA INC. [CA/CA]; 16711 Trans-Canada Highway, Kirkland, Quebec H9H 3L1 (CA).			
(72) Inventors; and (75) Inventors/Applicants (for US only): DUCHARME, Yves [CA/CA]; 4501 Kensington, Montreal, Quebec H4B 2W6 (CA). GAUTHIER, Jacques, Yves [CA/CA]; Apartment 2, 540 Odette Oigny, Laval, Quebec H7N 5Z4 (CA). PRASIT, Petpiboon [CA/CA]; 177 Argyle Drive, Kirkland, Quebec H9H 5A6 (CA). LEBLANC, Yves [CA/CA]; 8 Lafford, Kirkland, Quebec H9J 3Y3 (CA). WANG, Zhaoyin			Published Without international search report and to be republished upon receipt of that report.

(54) Title: PHENYL HETEROCYCLES AS CYCLOOXYGENASE-2 INHIBITORS

(57) Abstract

The invention encompasses the novel compound of formula (I) useful in the treatment of cyclooxygenase-2 mediated diseases. The invention also encompasses certain pharmaceutical compositions for treatment of cyclooxygenase-2 mediated diseases comprising compounds of formula (I).



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PHENYL HETEROCYCLES AS CYCLOOXYGENASE-2 INHIBITORS

BACKGROUND OF THE INVENTION

5 This invention relates to compounds and pharmaceutical compositions for the treatment of cyclooxygenase mediated diseases and methods of treatment thereof.

10 Non-steroidal, antiinflammatory drugs exert most of their antiinflammatory, analgesic and antipyretic activity and inhibit hormone-induced uterine contractions and certain types of cancer growth through inhibition of prostaglandin G/H synthase, also known as cyclooxygenase. Up until recently, only one form of cyclooxygenase had been characterized, this corresponding to cyclooxygenase-1 or the constitutive enzyme, as originally identified in bovine seminal vesicles. Recently the 15 gene for a second inducible form of cyclooxygenase (cyclooxygenase-2) has been cloned, sequenced and characterized from chicken, murine and human sources. This enzyme is distinct from the cyclooxygenase-1 which has now also been cloned, sequenced and characterized from sheep, murine and human sources. The second form of cyclooxygenase, cyclooxygenase-2, is rapidly and readily inducible by a number of agents including 20 mitogens, endotoxin, hormones, cytokines and growth factors. As prostaglandins have both physiological and pathological roles, we have concluded that the constitutive enzyme, cyclooxygenase-1, is responsible, in large part, for endogenous basal release of prostaglandins and hence is important in their physiological functions such as the maintenance of 25 gastrointestinal integrity and renal blood flow. In contrast, we have concluded that the inducible form, cyclooxygenase-2, is mainly responsible for the pathological effects of prostaglandins where rapid induction of the enzyme would occur in response to such agents as inflammatory agents, hormones, growth factors, and cytokines. Thus, a selective inhibitor of 30 cyclooxygenase-2 will have similar antiinflammatory, antipyretic and analgesic properties to a conventional non-steroidal antiinflammatory drug.

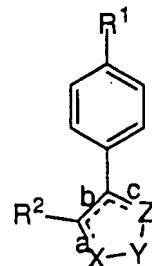
- 2 -

and in addition would inhibit hormone-induced uterine contractions and have potential anti-cancer effects, but will have a diminished ability to induce some of the mechanism-based side effects. In particular, such a compound should have a reduced potential for gastrointestinal toxicity, a reduced potential for renal side effects, a reduced effect on bleeding times 5 and possibly a lessened ability to induce asthma attacks in aspirin-sensitive asthmatic subjects.

SUMMARY OF THE INVENTION

10 The invention encompasses novel compounds of Formula I useful in the treatment of cyclooxygenase-2 mediated diseases.

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The invention also encompasses certain pharmaceutical compositions and methods for treatment of cyclooxygenase-2 mediated diseases comprising the use of compounds of Formula I.

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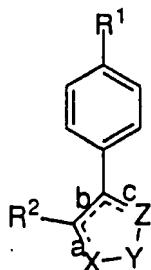
- 3 -

DETAILED DESCRIPTION OF THE INVENTION

The invention encompasses the novel compound of Formula I
useful in the treatment of cyclooxygenase-2 mediated diseases

5

10



I

or pharmaceutically acceptable salts thereof wherein:
X-Y-Z-is selected from the group consisting of:

15

- (a) -CH₂CH₂CH₂-,
- (b) -C(O)CH₂CH₂-,
- (c) -CH₂CH₂C(O)-,
- (d) -CR⁵(R^{5'})-O-C(O)-,
- (e) -C(O)-O-CR⁵(R^{5'})-,

20

- (f) -CH₂-NR³-CH₂-,
- (g) -CR⁵(R^{5'})-NR³-C(O)-,
- (h) -CR⁴=CR^{4'}-S-,
- (i) -S-CR⁴=CR^{4'}-,
- (j) -S-N=CH-,

25

- (k) -CH=N-S-,
- (l) -N=CR⁴-O-,
- (m) -O-CR⁴=N-
- (n) -N=CR⁴-NH-;
- (o) -N=CR⁴-S-, and

30

- (p) -S-CR⁴=N-;
- (q) -C(O)-NR³-CR⁵(R^{5'})-;

- 4 -

- (r) $-R^3N-CH=CH-$ provided R^1 is not $-S(O)2Me$
- (s) $-CH=CH-NR^3-$ provided R^1 is not $-S(O)2Me$

when side b is a double bond, and sides a and c are single bonds; and

5 $X-Y-Z$ is selected from the group consisting of:

- (a) $=CH-O-CH=$, and
- (b) $=CH-NR^3-CH=$,
- (c) $=N-S-CH=$,
- (d) $=CH-S-N=$,
- 10 (e) $=N-O-CH=$,
- (f) $=CH-O-N=$,
- (g) $=N-S-N=$,
- (h) $=N-O-N=$,

when sides a and c are double bonds and side b is a single bond;

15 R^1 is selected from the group consisting of

- (a) $S(O)2CH_3$,
- (b) $S(O)2NH_2$,
- (c) $S(O)2NHC(O)CF_3$,
- (d) $S(O)(NH)CH_3$,
- 20 (e) $S(O)(NH)NH_2$,
- (f) $S(O)(NH)NHC(O)CF_3$,
- (g) $P(O)(CH_3)OH$, and
- (h) $P(O)(CH_3)NH_2$,

R^2 is selected from the group consisting of

- (a) C_1 -6alkyl,
- (b) C_3, C_4, C_5, C_6 , and C_7 , cycloalkyl,
- (c) mono-, di- or tri-substituted phenyl or naphthyl wherein the substituent is selected from the group consisting of
 - (1) hydrogen,
 - (2) halo,
 - (3) C_1 -6alkoxy,

- 5 -

- (4) C₁₋₆alkylthio,
- (5) CN,
- (6) CF₃,
- (7) C₁₋₆alkyl,
- (8) N₃,
- 5 (9) -CO₂H,
- (10) -CO₂-C₁₋₄alkyl,
- (11) -C(R⁵)(R⁶)-OH,
- (12) -C(R⁵)(R⁶)-O-C₁₋₄alkyl, and
- (13) -C₁₋₆alkyl-CO₂-R⁵;
- 10 (d) mono-, di- or tri-substituted heteroaryl wherein the heteroaryl is a monocyclic aromatic ring of 5 atoms, said ring having one hetero atom which is S, O, or N, and optionally 1, 2, or 3 additionally N atoms; or
- 15 the heteroaryl is a monocyclic ring of 6 atoms, said ring having one hetero atom which is N, and optionally 1, 2, 3, or 4 additional N atoms; said substituents are selected from the group consisting of
 - (1) hydrogen,
 - (2) halo, including fluoro, chloro, bromo and iodo,
 - (3) C₁₋₆alkyl,
 - (4) C₁₋₆alkoxy,
 - (5) C₁₋₆alkylthio,
 - (6) CN,
 - (7) CF₃,
 - 25 (8) N₃,
 - (9) -C(R⁵)(R⁶)-OH, and
 - (10) -C(R⁵)(R⁶)-O-C₁₋₄alkyl;
- 20 (e) benzoheteroaryl which includes the benzo fused analogs of (d); R³ is selected from the group consisting of
 - (a) hydrogen,
 - (b) CF₃,

- 6 -

- (c) CN,
- (d) C₁₋₆alkyl,
- (e) hydroxyC₁₋₆alkyl,
- (f) -C(O)-C₁₋₆alkyl,
- (g) optionally substituted
 - (1) -C₁₋₅ alkyl-Q,
 - (2) -C₁₋₃alkyl-O-C₁₋₃alkyl-Q,
 - (3) -C₁₋₃alkyl-S-C₁₋₃alkyl-Q,
 - (4) -C₁₋₅ alkyl-O-Q, or
 - (5) -C₁₋₅ alkyl-S-Q,

10 wherein the substituent resides on the alkyl and the substituent is C₁₋₃alkyl;

- (h) -Q

R⁴ and R^{4'} are each independently selected from the group consisting of

- (a) hydrogen,
- (b) CF₃,
- (c) CN,
- (d) C₁₋₆alkyl,
- (e) -Q,
- (f) -O-Q;
- 20 (g) -S-Q, and
- (h) optionally substituted
 - (1) -C₁₋₅ alkyl-Q,
 - (2) -O-C₁₋₅ alkyl-Q,
 - (3) -S-C₁₋₅ alkyl-Q,
 - (4) -C₁₋₃alkyl-O-C₁₋₃alkyl-Q,
 - (5) -C₁₋₃alkyl-S-C₁₋₃alkyl-Q,
 - (6) -C₁₋₅ alkyl-O-Q,
 - (7) -C₁₋₅ alkyl-S-Q,

25 wherein the substituent resides on the alkyl and the substituent is C₁₋₃alkyl, and

- 7 -

R⁵, R^{5'}, R⁶, R⁷ and R⁸ are each independently selected from the group consisting of

- (a) hydrogen,
- (b) C₁-6alkyl,

5 or R⁵ and R⁶ or R⁷ and R⁸ together with the carbon to which they are attached form a saturated monocyclic carbon ring of 3, 4, 5, 6 or 7 atoms;

10 Q is CO₂H, CO₂-C₁-4alkyl, tetrazolyl-5-yl, C(R⁷)(R⁸)(OH), or

C(R⁷)(R⁸)(O-C₁-4alkyl);

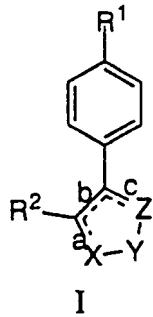
15

provided that when X-Y-Z is -S-CR⁴=CR^{4'}, then R⁴ and R^{4'} are other than CF₃.

15

In one aspect, within this embodiment are the compounds of formula I

20



or pharmaceutically acceptable salts thereof wherein:

25 X-Y-Z- is selected from the group consisting of -C(O)-O-CR⁵(R^{5'})- when side b is a double bond, and sides a and c are single bonds; and

R¹ is selected from the group consisting of

- (a) S(O)₂CH₃,
- (b) S(O)₂NH₂,

30

R² is selected from the group consisting of

- (a) C₁-6alkyl,
- (b) C₃, C₄, C₅, C₆, and C₇, cycloalkyl,

- 8 -

- (c) heteroaryl
- (d) benzoheteroaryl
- (e) mono- or di-substituted phenyl wherein the substituent is selected from the group consisting of
 - (1) hydrogen,
 - (2) halo,
 - (3) C₁₋₆alkoxy,
 - (4) C₁₋₆alkylthio,
 - (5) CN,
 - (6) CF₃,
 - (7) C₁₋₆alkyl,
 - (8) N₃,
 - (9) -CO₂H,
 - (10) -CO₂-C₁₋₄alkyl,
 - (11) -C(R⁵)(R⁶)-OH,
 - (12) -C(R⁵)(R⁶)-O-C₁₋₄alkyl, and
 - (13) -C₁₋₆alkyl-CO₂-R⁵;

R⁵, R^{5'} and R⁶ are each independently selected from the group consisting of

- (a) hydrogen,
- (b) C₁₋₆alkyl,

or R⁵ and R⁶ together with the carbon to which they are attached form a saturated monocyclic carbon ring of 3, 4, 5, 6 or 7 atoms.

One genus within the embodiment described above is the compound of formula I wherein

X-Y-Z-is selected from the group consisting of:

- (a) -CH₂CH₂CH₂-,
- (b) -C(O)CH₂CH₂-,
- (c) -CH₂CH₂C(O)-,
- (d) -CR⁵(R^{5'})-O-C(O)-,

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- (e) -C(O)-O-CR⁵(R^{5'})-,
- (f) -CH₂-NR³-CH₂-,
- (g) -CR⁵(R^{5'})-NR³-C(O)-,
- (h) -CR⁴=CR^{4'}-S-,
5 (i) -S-CR⁴=CR^{4'}-,
- (j) -S-N=CH-,
- (k) -CH=N-S-,
- (l) -N=CR⁴-O-,
- (m) -O-CR⁴=N-
- (n) -N-CR⁴-NH-,
10 (o) -N=CR⁴-S-, and
- (p) -S-CR⁴=N-,
- (q) -C(O)-NR³-CR⁵(R^{5'})-;
- (r) -NR³-CH=CH- provided R¹ is other than -S(O)₂Me,
- (s) -CH=CH-NR³- provided R¹ is other than -S(O)₂Me.
15

Within this genus is the sub-genus of compounds of formula I
wherein

R¹ is selected from the group consisting of

- (a) S(O)₂CH₃,
- 20 (b) S(O)₂NH₂,
- (c) S(O)₂NHC(O)CF₃,
- (d) S(O)NHCH₃,
- (e) S(O)NHNH₂, and
- (f) S(O)NHNHC(O)CF₃;

25 R² is selected from the group consisting of

- (a) C₁-4alkyl,
- (b) C₃, C₄, C₅, C₆, and C₇, cycloalkyl,
- (c) mono- or di-substituted phenyl wherein the substituent is
30 selected from the group consisting of
 - (1) hydrogen,
 - (2) fluoro, chloro, and bromo,

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- (3) C₁-4alkoxy,
- (4) C₁-4alkylthio,
- (5) CN,
- (6) CF₃,
- (7) C₁-4alkyl,
- 5 (8) N₃,
- (9) -CO₂H,
- (10) -CO₂-C₁-3alkyl,
- (11) -C(R⁵)(R⁶)-OH, and
- (12) -C(R⁵)(R⁶)-O-C₁-3alkyl,
- 10 (d) mono- or di-substituted heteroaryl selected from the group consisting of
 - (1) furanyl,
 - (2) diazinyl, triazinyl and tetrazinyl,
 - (3) imidazolyl,
 - 15 (4) isooxazolyl,
 - (5) isothiazolyl,
 - (6) oxadiazolyl,
 - (7) oxazolyl,
 - (8) pyrazolyl,
 - 20 (9) pyrrolyl,
 - (10) thiadiazolyl,
 - (11) thiazolyl,
 - (12) thienyl,
 - (13) triazolyl, and
 - 25 (14) tetrazolyl,
- wherein said substituents are selected from the group consisting of
 - (a) hydrogen,
 - (b) fluoro, chloro, bromo,
 - (c) C₁-4alkoxy,
 - 30 (d) C₁-4alkylthio,
 - (e) CN,

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- (5) CF₃,
- (6) C₁₋₄alkyl,
- (7) N₃,
- (8) -C(R⁵)(R⁶)-OH,
- (9) -C(R⁵)(R⁶)-O-C₁₋₄alkyl.

5

Within this sub-genus is the class of compounds of formula I
wherein

R² is selected from the group consisting of

- (a) cyclohexyl, and
- 10 (b) mono- or di-substituted phenyl, and
wherein the substitutents are selected from the group
consisting of
 - (1) hydrogen,
 - (2) halo,
 - 15 (3) C₁₋₄alkoxy,
 - (4) C₁₋₄alkylthio,
 - (5) CN,
 - (6) CF₃,
 - (7) C₁₋₄alkyl,
 - 20 (8) N₃, and
 - (9) -C(R⁵)(R⁶)-OH;

R³ is selected from the group consisting of

- (a) hydrogen,
- (b) CF₃,
- 25 (c) C₁₋₃alkyl and hydroxyC₁₋₃alkyl,
- (d) CN,

R⁴ and R^{4'} are each independently selected from the group consisting of

- (a) hydrogen,
- (b) CF₃,
- 30 (c) C₁₋₃alkyl,
- (d) CN,

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(e) chloro and fluoro; and

R⁵, R^{5'}, R⁶, are each independently selected from the group consisting of

(a) hydrogen,

(b) methyl or ethyl,

or R⁵ and R⁶ together with the carbon to which they are attached

5 form a saturated carbon ring of 4, 5 or 6 atoms,

Within this class is the sub-class of compounds of formula I wherein X-Y-Z-is selected from the group consisting of:

(a) -CH₂-O-C(O)-,

10 (b) -C(O)-O-CH₂-, and

(c) -CH₂-NR³-C(O)-;

R¹ is selected from the group consisting of

(a) S(O)₂CH₃,

(b) S(O)₂NH₂,

15 (c) S(O)NHCH₃, and

(d) S(O)NHNH₂;

R² is selected from the group consisting of

mono or di-substituted phenyl wherein the substitutents are selected from the group consisting of

20 (1) hydrogen,

(2) halo, selected from the group consisting of fluoro, chloro and bromo,

(3) C₁-3alkoxy,

(4) C₁-3alkylthio,

25 (5) CN, and

(6) C₁-3alkyl;

R³ is selected from the group consisting of

(a) hydrogen,

(b) CF₃,

30 (c) C₁-3alkyl and hydroxyC₁-3alkyl.

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Within this sub-class is the group of compounds of formula I
wherein

X-Y-Z-is selected from the group consisting of:

- (a) -CH₂-O-C(O)-, and
- (b) -C(O)-O-CH₂-, and

5 R¹ is selected from the group consisting of

- (a) S(O)₂CH₃,
- (b) S(O)₂NH₂,
- (c) S(O)NHCH₃, and
- (d) S(O)NHNH₂;

10 R² is

mono or di-substituted phenyl wherein the substitutents are selected from the group consisting of

- (1) hydrogen,
- (2) halo, selected from the group consisting of fluoro, chloro and bromo,
- (3) methoxy, and
- (4) methyl.

20 This group may be more particularly defined as the compounds of formula I wherein

X-Y-Z-is selected from the group consisting of:

- (a) -CH₂-O-C(O)-, and
- (b) -C(O)-O-CH₂-, and

25 R¹ is selected from the group consisting of

- (a) S(O)₂CH₃, and
- (b) S(O)₂NH₂,

R² is

mono or di-substituted phenyl wherein the substitutents are selected from the group consisting of

30 (1) hydrogen,

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(2) halo, selected from the group consisting of fluoro, chloro and bromo.

Within the sub-genus escribed above there is the class of compounds of formula I wherein

5 R² is a mono- or di-substituted heteroaryl wherein heteroaryl is selected from the group consisting of

(1) furanyl,
(2) diazinyl, triazinyl, tetrazinyl,
(3) imidazolyl,
10 (4) isooxazolyl,
(5) isothiazolyl,
(6) oxadiazolyl,
(7) oxazolyl,
(8) pyrazolyl,
15 (9) pyrrolyl,
(10) thiadiazolyl,
(11) thiazolyl,
(12) thienyl,
(13) triazolyl, and
20 (14) tetrazolyl,

wherein the substitutents are selected from the group consisting of

(a) hydrogen,
(b) fluoro or chloro,
25 (c) C₁₋₃alkoxy,
(d) C₁₋₆alkylthio,
(e) CN,
(f) CF₃,
(g) C₁₋₃alkyl,
30 (h) -C(R⁵)(R⁶)-OH;
(i) -C(R⁵)(R⁶)-O-C₁₋₄alkyl.

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Within this class there is the sub-class of compounds of formula I wherein

R^2 is a mono- or di-substituted heteroaryl wherein heteroaryl is selected from the group consisting of

- 5 (1) 2-furanyl,
- (2) 3-furanyl,
- (3) 2-thienyl,
- (4) 3-thienyl,
- (5) 3-isoxazolyl,
- 10 (6) 4-isoxazolyl,
- (7) 5-isoxazolyl,
- (8) 3-isothiazolyl,
- (9) 4-isothiazolyl,
- (10) 5-isothiazolyl,
- 15 (11) 2-oxazolyl,
- (12) 4-oxazolyl,
- (13) 5-oxazolyl,
- (14) 2-thiazolyl,
- (15) 4-thiazolyl,
- 20 (16) 5-thiazolyl,
- (17) 1,2,3-thiadiazol-4-yl,
- (18) 1,2,3-thiadiazol-5-yl,
- (19) 1,2,4-thiadiazol-3-yl,
- (20) 1,2,4-thiadiazol-5-yl,
- 25 (21) 1,3,4-thiadiazol-2-yl,
- (22) 1,2,5-thiadiazol-3-yl,
- (23) 1,2,3-oxadiazol-4-yl,
- (24) 1,2,3-oxadiazol-5-yl,
- (25) 1,2,4-oxadiazol-3-yl,
- 30 (26) 1,2,4-oxadiazol-5-yl,
- (27) 1,3,4-oxadiazol-2-yl,

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- (28) 1,2,5-oxadiazol-3-yl,
- (29) pyrazol-4-yl,
- (30) pyrazol-5-yl,
- (31) 1,2,3-triazol-4-yl,
- (32) 1,2,3-triazol-5-yl,
- (33) 1,2,4-triazol-3-yl,
- (34) 1,2,4-triazol-5-yl,
- (35) 1,2-diazinyl,
- (36) 1,3-diazinyl,
- (37) 1,4-diazinyl,
- (38) 1,2,3,4-tetrazin-5-yl,
- (39) 1,2,4,5-tetrazin-4-yl,
- (40) 1,3,4,5-tetrazin-2-yl, and
- (41) 1,2,3,5-tetrazin-4-yl.

15 Within this sub-class there is the group of compounds of formula I wherein the heteroaryl is selected from the group consisting of

- (1) 3-isoxazolyl,
- (2) 4-isoxazolyl,
- (3) 5-isoxazolyl,
- (4) 3-isothiazolyl,
- (5) 4-isothiazolyl,
- (6) 5-isothiazolyl,
- (7) 2-oxazolyl,
- (8) 4-oxazolyl,
- (9) 5-oxazolyl,
- (10) 2-thiazolyl,
- (11) 4-thiazolyl,
- (12) 5-thiazolyl,
- (13) 1,2,3-thiadiazol-4-yl,
- (14) 1,2,3-thiadiazol-5-yl,
- (15) 1,2,4-thiadiazol-3-yl,

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- (16) 1,2,4-thiadiazol-5-yl,
- (17) 1,3,4-thiadiazol-2-yl,
- (18) 1,2,5-thiadiazol-3-yl,
- (19) 1,2,3-oxadiazol-4-yl,
- 5 (20) 1,2,3-oxadiazol-5-yl,
- (21) 1,2,4-oxadiazol-3-yl,
- (22) 1,2,4-oxadiazol-5-yl,
- (23) 1,3,4-oxadiazol-2-yl,
- (24) 1,2,5-oxadiazol-3-yl,
- (25) 1,2-diazinyl,
- 10 (26) 1,3-diazinyl, and
- (27) 1,4-diazinyl.

These heteroaryls may be more particularly defined as being selected from the group consisting of

- 15 (1) 3-isothiazolyl,
- (2) 4-isothiazolyl,
- (3) 5-isothiazolyl,
- (4) 2-oxazolyl,
- (5) 4-oxazolyl,
- 20 (6) 5-oxazolyl,
- (7) 2-thiazolyl,
- (8) 4-thiazolyl,
- (9) 5-thiazolyl,
- (10) 1,2-diazinyl,
- 25 (11) 1,3-diazinyl, and
- (12) 1,4-diazinyl, and

wherein the substituents are selected from the group consisting of

- (1) hydrogen,
- (2) fluoro or chloro,
- 30 (3) C₁₋₃alkoxy,
- (4) C₁₋₃alkylthio,

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(5) CN,

(6) C₁-3alkyl, and

(7) -C(R⁵)(R⁶)-OH,

wherein R⁵ and R⁶ are each independently hydrogen, methyl or ethyl.

5

and may be further particularly

Given these more particularly defined definitions of heteroaryl, the compounds of formula I includes the group wherein

X-Y-Z-is selected from the group consisting of:

10

(a) -CH₂-O-C(O)-,

(b) -C(O)-O-CH₂-, and

(c) -CH₂-NR³-C(O)-;

R¹ is selected from the group consisting of

15

(a) S(O)₂CH₃,

(b) S(O)₂NH₂,

(c) S(O)NHCH₃, and

(d) S(O)NHNH₂, and

R³ is selected from the group consisting of

20

(a) hydrogen,

(b) CF₃,

(c) C₁-3alkyl and hydroxyC₁-3alkyl,

(d) CN.

25 A second genus within the embodiment described above is the compounds of formula I wherein

X-Y-Z-is selected from the group consisting of:

30

(a) =CH-O-CH=, and

(b) =CH-NR³-CH=,

(c) =N-S-CH=,

(d) =CH-S-N=,

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- (e) =N-O-CH=,
- (f) =CH-O-N=,
- (g) =N-S-N=,
- (h) =N-O-N=.

5 Within this genus is the sub-genus of compounds of formula I
wherein

R¹ is selected from the group consisting of

- (a) S(O)2CH₃,
- (b) S(O)2NH₂,
- (c) S(O)2NHC(O)CF₃,
- (d) S(O)(NH)CH₃,
- (e) S(O)(NH)NH₂, and
- (f) S(O)(NH)NHC(O)CF₃;

R² is selected from the group consisting of

- (a) C₁₋₄alkyl,
- (b) C₃, C₄, C₅, C₆, and C₇, cycloalkyl,
- (c) mono- or di-substituted phenyl wherein the substituent is selected from the group consisting of
 - (1) hydrogen,
 - (2) fluoro, chloro, and bromo,
 - (3) C₁₋₄alkoxy,
 - (4) C₁₋₄alkylthio,
 - (5) CN,
 - (6) CF₃,
 - (7) C₁₋₄alkyl,
 - (8) N₃,
 - (9) -CO₂H,
 - (10) -CO₂-C₁₋₃alkyl,
 - (10) -C(R⁵)(R⁶)-OH, and
 - (11) -C(R⁵)(R⁶)-O-C₁₋₃alkyl,

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(d) mono- or di-substituted heteroaryl selected from the group consisting of

5 (1) furanyl,
(2) diazinyl, triazinyl and tetrazinyl,
(3) imidazolyl,
(4) isooxazolyl,
(5) isothiazolyl,
10 (6) oxadiazolyl,
(7) oxazolyl,
(8) pyrazolyl,
(9) pyrrolyl,
(10) thiadiazolyl,
(11) thiazolyl,
(12) thienyl,
15 (13) triazolyl, and
(14) tetrazolyl,

wherein said substituents are selected from the group consisting of

20 (a) hydrogen,
(b) fluoro, chloro, bromo,
(c) C₁-4alkoxy,
(d) C₁-4alkylthio,
(e) CN,
(5) CF₃,
(6) C₁-4alkyl,
(7) N₃,
25 (8) -C(R⁵)(R⁶)-OH;
(9) -C(R⁵)(R⁶)-O-C₁-4alkyl.

For purposes of this specification the heteroaryls of this sub-genus may be more particularly described in any of the manners described above.

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Within this sub-genus there is the class of compounds of formula I wherein

R² is selected from the group consisting of

- (a) cyclohexyl, and
- (b) mono or di substituted phenyl, and

5 wherein the substitutents are selected from the group consisting of

(1) hydrogen,

(2) halo,

(3) C₁₋₄alkoxy,

10 (4) C₁₋₄alkylthio,

(5) CN,

(6) CF₃,

(7) C₁₋₄alkyl,

(8) N₃, and

15 (9) -C(R⁵)(R⁶)-OH;

R³ is selected from the group consisting of

- (a) hydrogen,
- (b) CF₃,
- (c) C₁₋₃alkyl and hydroxyC₁₋₃alkyl,
- (d) CN;

R⁵, R^{5'}, R⁶, are each independently selected from the group consisting of

- (a) hydrogen,
- (b) methyl or ethyl,

or R⁵ and R⁶ together with the carbon to which they are attached
25 form a saturated carbon ring of 4, 5 or 6 atoms.

Within this class there is the sub-class of compounds of formula I wherein

X-Y-Z-is selected from the group consisting of:

- 30 (a) =CH-O-CH=,
- (b) =N-S-N=,

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(c) =N-O-N=;

R¹ is selected from the group consisting of

- (a) S(O)2CH₃, and
- (b) S(O)2NH₂;

R² is selected from the group consisting of

5 mono- or di-substituted phenyl wherein the substituents are selected from the group consisting of

- (1) hydrogen,
- (2) halo, selected from the group consisting of fluoro, chloro and bromo,
- 10 (3) C₁-3alkoxy,
- (4) C₁-3alkylthio,
- (5) CF₃,
- (6) C₁-3alkyl;

R³ is selected from the group consisting of

15 (a) hydrogen,

- (b) CF₃,
- (c) C₁-3alkyl and hydroxyC₁-3alkyl,

R⁵ and R⁶ are each selected from the group consisting of

20 (a) hydrogen,

- (b) methyl or ethyl,

or R⁵, R^{5'} and R⁶ together with the carbon to which they are attached form a saturated carbon ring of 5, 6 or 7 atoms.

For purposes of this specification alkyl is defined to include

25 linear, branched, and cyclic structures, with C₁-6alkyl including methyl, ethyl, propyl, 2-propyl, s- and t-butyl, butyl, pentyl, hexyl, 1,1-dimethylethyl, cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl. Similarly, C₁-6alkoxy is intended to include alkoxy groups of from 1 to 6 carbon atoms of a straight, branched, or cyclic configuration. Examples of 30 lower alkoxy groups include methoxy, ethoxy, propoxy, isopropoxy, cyclopropoxy, cyclohexyloxy, and the like. Likewise, C₁-6alkylthio is

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intended to include alkylthio groups of from 1 to 6 carbon atoms of a straight, branched or cyclic configuration. Examples of lower alkylthio groups include methylthio, propylthio, isopropylthio, cycloheptylthio, etc. By way of illustration, the propylthio group signifies -SCH₂CH₂CH₃.

5 Heteroaryl includes furan, thiophene, pyrrole, isoxazole, isothiazole, pyrazole, oxazole, thiazole, imidazole, 1,2,3-oxadiazole, 1,2,3-thiadiazole, 1,2,3-triazole, 1,3,4-oxadiazole, 1,3,4-thiadiazole, 1,3,4-triazole, 1,2,5-oxadiazole, 1,2,5-thiadiazole, pyridine, pyridazine, pyrimidine, pyrazine, 1,2,4-triazine, 1,3,5-triazine, 1,2,4,5-tetrazine, and the like.

10 Benzoheteroaryl includes the above heteroaryl rings to which it is possible to fuse a benzene ring.

Exemplifying the invention are:

- (a) 3-(4-(Aminosulfonyl)phenyl)-2-(4-fluorophenyl)-5-(2-hydroxy-2-propyl)thiophene,
- (b) 3-(4-(Aminosulfonyl)phenyl)-2-(4-fluorophenyl)thiophene,
- (c) 3-(4-(Aminosulfonyl)phenyl)-2-(4-fluorophenyl)-5-(2-propyl)thiophene,
- (d) 3-(4-(Aminosulfonyl)phenyl)-2-cyclohexylthiophene,
- (e) 5-(4-Carboxyphenyl)-4-(4-(methylsulfonyl)phenyl)thiophene-2-carboxylic acid,
- (f) 4-(4-Fluorophenyl)-2-methyl-5-(4-(methylsulfonyl)phenyl)thiazole,
- (g) 2-(4-Fluorophenyl)-3-(4-(methylsulfonyl)phenyl)-2-cyclopenten-1-one
- (h) 4-(4-(Methylsulfonyl)phenyl)-5-(4-fluorophenyl)-isothiazole,
- (i) 3-(4-Fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone,
- (j) 3-(4-Fluorophenyl)-4-(4-(aminosulfonyl)phenyl)-2-(5H)-furanone,
- (k) 3-(4-Fluorophenyl)-4-(4-(methylsulfonyl)phenyl)furan,

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- (l) 5,5-Dimethyl-3-(4-fluorophenyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (m) 2-(4-(Aminosulfonyl)phenyl)-3-(4-fluorophenyl)thiophene, and
- 5 (n) 3-(4-(Trifluoroacetylaminosulfonyl)phenyl)-2-(4-fluorophenyl)thiophene,
- (o) 3-(3-Fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (p) 5,5-Dimethyl-3-(3-fluorophenyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- 10 (q) 5,5-Dimethyl-3-(3-chlorophenyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (r) 3-(3,4-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (s) 3-(3,4-Dichlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- 15 (t) 5,5-Dimethyl-3-(3,4-difluorophenyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (u) 5,5-Dimethyl-3-(3,4-dichlorophenyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- 20 (v) 5,5-Dimethyl-3-(4-chlorophenyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (w) 3-(2-Naphyhyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (x) 5,5-Dimethyl-3-(2-naphyhyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- 25 (y) 3-phenyl-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone.

Further illustrating the invention are

- (a) 3-(3,4-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone, and

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(b) 3-phenyl-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone, or a pharmaceutically acceptable salt thereof.

Some of the compounds described herein contain one or more asymmetric centers and may thus give rise to diastereomers and optical isomers. The present invention is meant to comprehend such possible diastereomers as well as their racemic and resolved, enantiomerically pure forms and pharmaceutically acceptable salts thereof.

Some of the compounds described herein contain olefinic double bonds, and unless specified otherwise, are meant to include both E and Z geometric isomers.

In a second embodiment, the invention encompasses pharmaceutical compositions for inhibiting cyclooxygenase and for treating cyclooxygenase mediated diseases as disclosed herein comprising a pharmaceutically acceptable carrier and a non-toxic therapeutically effective amount of compound of formula I as described above.

Within this embodiment the invention encompasses pharmaceutical compositions for inhibiting cyclooxygenase-2 and for treating cyclooxygenase-2 mediated diseases as disclosed herein comprising a pharmaceutically acceptable carrier and a non-toxic therapeutically effective amount of compound of formula I as described above.

In a third embodiment, the invention encompasses a method of inhibiting cyclooxygenase and treating cyclooxygenase mediated diseases, advantageously treated by an active agent that selectively inhibits COX-2 in preference to COX-1 as disclosed herein comprising: administration to a patient in need of such treatment of a non-toxic therapeutically effective amount of a compound of Formula I as disclosed herein.

For purposes of this specification a compound is said to selectively inhibit COX-2 in preference to COX-1 if the ratio of the IC50 concentration for COX-1 inhibition to COX-2 inhibition is 100 or greater.

The pharmaceutical compositions of the present invention comprise a compound of Formula I as an active ingredient or a pharmaceutically acceptable salt, thereof, and may also contain a pharmaceutically acceptable carrier and optionally other therapeutic ingredients. The term "pharmaceutically acceptable salts" refers to salts prepared from pharmaceutically acceptable non-toxic bases including inorganic bases and organic bases. Salts derived from inorganic bases include aluminum, ammonium, calcium, copper, ferric, ferrous, lithium, magnesium, manganic salts, manganous, potassium, sodium, zinc, and the like. Particularly preferred are the ammonium, calcium, magnesium, potassium, and sodium salts. Salts derived from pharmaceutically acceptable organic non-toxic bases include salts of primary, secondary, and tertiary amines, substituted amines including naturally occurring substituted amines, cyclic amines, and basic ion exchange resins, such as arginine, betaine, caffeine, choline, N,N--dibenzylethylenediamine, diethylamine, 2-diethylaminoethanol, 2-dimethylaminoethanol, ethanolamine, ethylenediamine, N-ethylmorpholine, N-ethylpiperidine, glucamine, glucosamine, histidine, hydrabamine, isopropylamine, lysine, methylglucamine, morpholine, piperazine, piperidine, polyamine resins, procaine, purines, theobromine, triethylamine, trimethylamine, tripropylamine, tromethamine, and the like.

It will be understood that in the discussion of methods of treatment which follows, references to the compounds of Formula I are meant to also include the pharmaceutically acceptable salts.

The Compound of Formula I is useful for the relief of pain, fever and inflammation of a variety of conditions including rheumatic fever, symptoms associated with influenza or other viral infections, common cold, low back and neck pain, dysmenorrhea, headache, toothache, sprains and strains, myositis, neuralgia, synovitis, arthritis, including rheumatoid arthritis degenerative joint diseases (osteoarthritis), gout and ankylosing spondylitis, bursitis, burns, injuries, following surgical and dental procedures. In addition, such a compound may inhibit cellular

neoplastic transformations and metastatic tumor growth and hence can be used in the treatment of cancer. Compounds of formula I may also be useful for the treatment of dementia including pre-senile and senile dementia, and in particular, dementia associated with Alzheimer Disease (ie Alzheimer's dementia).

5 Compounds of formula I will also inhibit prostanoid-induced smooth muscle contraction by preventing the synthesis of contractile prostanoids and hence may be of use in the treatment of dysmenorrhea, premature labor and asthma.

10 By virtue of its high cyclooxygenase-2 (COX-2) activity and/or its selectivity for cyclooxygenase-2 over cyclooxygenase-1 (COX-1) as defined above, compounds of formula I will prove useful as an alternative to conventional non-steroidal antiinflammatory drugs (NSAID'S) particularly where such non-steroidal antiinflammatory drugs may be contra-indicated such as in patients with peptic ulcers, gastritis, 15 regional enteritis, ulcerative colitis, diverticulitis or with a recurrent history of gastrointestinal lesions; GI bleeding, coagulation disorders including anemia such as hypoprothrombinemia, haemophilia or other bleeding problems (including those relating to reduced or impaired platelet function); kidney disease (eg impaired renal function); those prior to 20 surgery or taking anticoagulants; and those susceptible to NSAID induced asthma.

25 Similarly, compounds of formula I, will be useful as a partial or complete substitute for conventional NSAID'S in preparations wherein they are presently co-administered with other agents or ingredients. Thus in further aspects, the invention encompasses pharmaceutical compositions for treating cyclooxygenase-2 mediated diseases as defined above comprising a non-toxic therapeutically effective amount of the compound of Formula I as defined above and one or more ingredients such as another pain reliever including acetominophen or phenacetin; a potentiator 30 including caffeine; an H2-antagonist, aluminum or magnesium hydroxide, simethicone, a decongestant including phenylephrine,

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phenylpropanolamine, pseudophedrine, oxymetazoline, ephinephrine, naphazoline, xylometazoline, propylhexedrine, or levo-desoxyephedrine; an antiitussive including codeine, hydrocodone, caramiphen, carbetapentane, or dextramethorphan; a diuretic; a sedating or non-sedating antihistamine. In addition the invention encompasses a method of treating 5 cyclooxygenase mediated diseases comprising: administration to a patient in need of such treatment a non-toxic therapeutically effect amount of the compound of Formula I, optionally co-administered with one or more of such ingredients as listed immediately above.

10 Compounds of the present invention are inhibitors of cyclooxygenase-2 and are thereby useful in the treatment of cyclooxygenase-2 mediated diseases as enumerated above. This activity is illustrated by their ability to selectively inhibit cyclooxygenase-2 over cyclooxygenase-1. Accordingly, in one assay, the ability of the compounds 15 of this invention to treat cyclooxygenase mediated diseases can be demonstrated by measuring the amount of prostaglandin E2 (PGE2) synthesized in the presence of arachidonic acid, cyclooxygenase-1 or cyclooxygenase-2 and a compound of formula I. The IC50 values represent the concentration of inhibitor required to return PGE2 synthesis 20 to 50 % of that obtained as compared to the uninhibited control. Illustrating this aspect, we have found that the Compounds of the Examples are more than 100 times more effective in inhibiting COX-2 than they are at inhibiting COX-1. In addition they all have a COX-2 IC50 of 1 nM to 1 μ M. By way of comparison, Ibuprofen has an IC50 for COX-2 of 1 μ M, 25 and Indomethacin has an IC50 for COX-2 of approximately 100 nM. For the treatment of any of these cyclooxygenase mediated diseases, compounds of formula I may be administered orally, topically, parenterally, by inhalation spray or rectally in dosage unit formulations containing conventional non-toxic pharmaceutically acceptable carriers, 30 adjuvants and vehicles. The term parenteral as used herein includes subcutaneous injections, intravenous, intramuscular, intrasternal injection

or infusion techniques. In addition to the treatment of warm-blooded animals such as mice, rats, horses, cattle sheep, dogs, cats, etc., the compound of the invention is effective in the treatment of humans.

As indicated above, pharmaceutical compositions for treating cyclooxygenase-2 mediated diseases as defined may optionally include one or more ingredients as listed above.

The pharmaceutical compositions containing the active ingredient may be in a form suitable for oral use, for example, as tablets, troches, lozenges, aqueous or oily suspensions, dispersible powders or granules, emulsions, hard or soft capsules, or syrups or elixirs.

10 Compositions intended for oral use may be prepared according to any method known to the art for the manufacture of pharmaceutical compositions and such compositions may contain one or more agents selected from the group consisting of sweetening agents, flavoring agents, coloring agents and preserving agents in order to provide pharmaceutically elegant and palatable preparations. Tablets contain the active ingredient in admixture with non-toxic pharmaceutically acceptable excipients which are suitable for the manufacture of tablets. These excipients may be for example, inert diluents, such as calcium carbonate, sodium carbonate, lactose, calcium phosphate or sodium phosphate; granulating and 15 disintegrating agents, for example, corn starch, or alginic acid; binding agents, for example starch, gelatin or acacia, and lubricating agents, for example, magnesium stearate, stearic acid or talc. The tablets may be uncoated or they may be coated by known techniques to delay 20 disintegration and absorption in the gastrointestinal tract and thereby provide a sustained action over a longer period. For example, a time delay material such as glyceryl monostearate or glyceryl distearate may be 25 employed. They may also be coated by the technique described in the U.S. Patent 4,256,108; 4,166,452; and 4,265,874 to form osmotic therapeutic tablets for control release.

30 Formulations for oral use may also be presented as hard gelatin capsules wherein the active ingredient is mixed with an inert solid

diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules wherein the active ingredients is mixed with water or an oil medium, for example peanut oil, liquid paraffin, or olive oil.

Aqueous suspensions contain the active material in admixture with excipients suitable for the manufacture of aqueous suspensions. Such 5 excipients are suspending agents, for example sodium carboxymethyl-cellulose, methylcellulose, hydroxy-propylmethycellulose, sodium alginate, polyvinyl-pyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents may be a naturally-occurring phosphatide, for example lecithin, or condensation products of an alkylene oxide with fatty acids, for 10 example polyoxyethylene stearate, or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethylene-oxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with partial esters 15 derived from fatty acids and hexitol anhydrides, for example polyethylene sorbitan monooleate. The aqueous suspensions may also contain one or more preservatives, for example ethyl, or n-propyl, p-hydroxybenzoate, one or more coloring agents, one or more flavoring agents, and one or more sweetening agents, such as sucrose, saccharin or aspartame.

20 Oily suspensions may be formulated by suspending the active ingredient in a vegetable oil, for example arachis oil, olive oil, sesame oil or coconut oil, or in mineral oil such as liquid paraffin. The oily suspensions may contain a thickening agent, for example beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set forth above, 25 and flavoring agents may be added to provide a palatable oral preparation. These compositions may be preserved by the addition of an anti-oxidant such as ascorbic acid.

30 Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water provide the active ingredient in admixture with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or wetting agents

and suspending agents are exemplified by those already mentioned above. Additional excipients, for example sweetening, flavoring and coloring agents, may also be present.

The pharmaceutical compositions of the invention may also be in the form of an oil-in-water emulsions. The oily phase may be a 5 vegetable oil, for example olive oil or arachis oil, or a mineral oil, for example liquid paraffin or mixtures of these. Suitable emulsifying agents may be naturally-occurring phosphatides, for example soy bean, lecithin, and esters or partial esters derived from fatty acids and hexitol anhydrides, for example sorbitan monooleate, and condensation products of the said 10 partial esters with ethylene oxide, for example polyoxyethylene sorbitan monooleate. The emulsions may also contain sweetening and flavouring agents.

Syrups and elixirs may be formulated with sweetening agents, for example glycerol, propylene glycol, sorbitol or sucrose. Such 15 formulations may also contain a demulcent, a preservative and flavoring and coloring agents. The pharmaceutical compositions may be in the form of a sterile injectable aqueous or oleagenous suspension. This suspension may be formulated according to the known art using those suitable dispersing or wetting agents and suspending agents which have been 20 mentioned above. The sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example as a solution in 1,3-butane diol. Among the acceptable vehicles and solvents that may be employed are water, Ringer's solution and isotonic sodium chloride solution. In addition, sterile, fixed 25 oils are conventionally employed as a solvent or suspending medium. For this purpose any bland fixed oil may be employed including synthetic mono- or diglycerides. In addition, fatty acids such as oleic acid find use in the preparation of injectables.

Compounds of formula I may also be administered in the form 30 of a suppositories for rectal administration of the drug. These compositions can be prepared by mixing the drug with a suitable non-

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irritating excipient which is solid at ordinary temperatures but liquid at the rectal temperature and will therefore melt in the rectum to release the drug. Such materials are cocoa butter and polyethylene glycols.

For topical use, creams, ointments, jellies, solutions or suspensions, etc., containing the compound of Formula I are employed. (For purposes of this application, topical application shall include mouth washes and gargles.)

Dosage levels of the order of from about 0.01 mg to about 140 mg/kg of body weight per day are useful in the treatment of the above-indicated conditions, or alternatively about 0.5 mg to about 7 g per patient per day. For example, inflammation may be effectively treated by the administration of from about 0.01 to 50 mg of the compound per kilogram of body weight per day, or alternatively about 0.5 mg to about 3.5 g per patient per day, preferably 2.5 mg to 1 g per patient per day.

The amount of active ingredient that may be combined with the carrier materials to produce a single dosage form will vary depending upon the host treated and the particular mode of administration. For example, a formulation intended for the oral administration of humans may contain from 0.5 mg to 5 g of active agent compounded with an appropriate and convenient amount of carrier material which may vary from about 5 to about 95 percent of the total composition. Dosage unit forms will generally contain between from about 1 mg to about 500 mg of an active ingredient, typically 25 mg, 50 mg, 100 mg, 200 mg, 300 mg, 400 mg, 500 mg, 600 mg, 800 mg, or 1000 mg.

It will be understood, however, that the specific dose level for any particular patient will depend upon a variety of factors including the age, body weight, general health, sex, diet, time of administration, route of administration, rate of excretion, drug combination and the severity of the particular disease undergoing therapy.

Methods of Synthesis

The compounds of the present invention can be prepared according to the following methods.

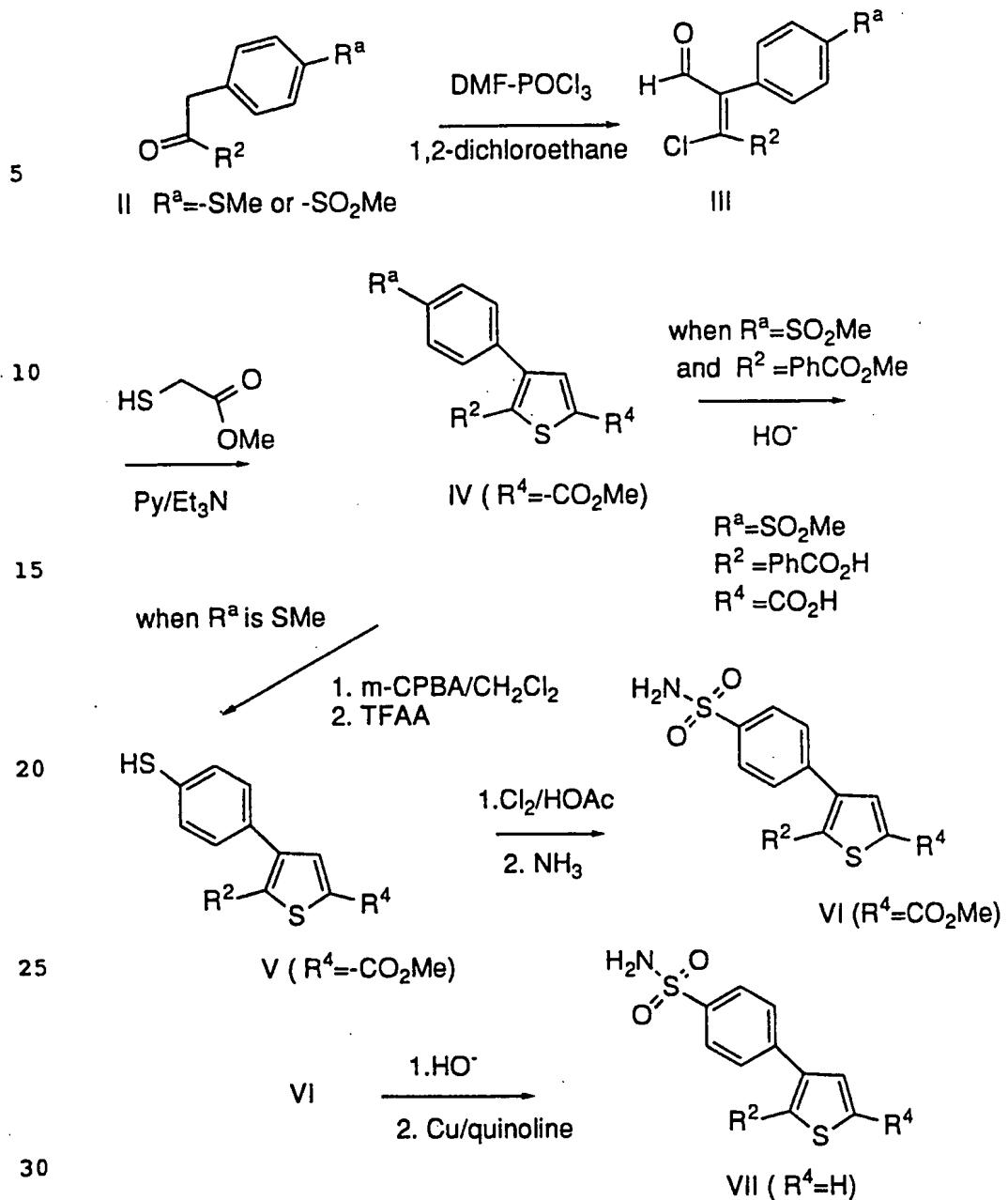
Method A:

The β -chlorovinylaldehyde III can be obtained from the 5 ketone II and the Vilsmeier reagent (DMF-POCl₃) using the general method described by Weissenfels (Z. Chem. 1966, 6, 471). The thiophene compound IV is obtained from III using the general method described by Weissenfels (Z. Chem., 1973, 13, 57). The thiol compound V can be obtained after oxidation of compound IV ($R^a = -SMe$) with one equivalent 10 of m-CPBA followed by treatment of the resulting sulfoxide with TFAA at reflux. The sulfonamide group (VI) can then be formed by the method of Kharash (J. Amer. Chem. Soc. 1951, 73, 3240). The hydrolysis of compound VI and decarboxylation with Cu bronze in quinoline provides compound VII. Compound VII ($R^4 = H$) can be treated with halogenating 15 agent such as bromine in acetic acid to allow the preparation of the 5-bromothiophene (VII, $R^4 = Br$). When it is desired to have a nitrile group at C-5, this can be accomplished from VI via amide formation using the Weinreb methodology (Tetrahedron Letters, 1977, 4171) followed by 20 dehydration with TFAA. The CF₃ group can be introduced at C-5 of VII via the method of Girard (J. Org. Chem. 1983, 48, 3220).

The introduction of an alkyl group at C-5 can be achieved via a Friedel-Crafts reaction on VII ($R^4 = H$) and an acyl chloride, Cl-CO-lower alkyl and a catalyst such as TiCl₄, followed by reduction. For $R^4 = Me$, this can be achieved from the ester ($R^4 = CO_2Me$) via a DIBAL-H reduction followed by deoxygenation using the method of Lau (J. Org. Chem. 1986, 51, 3038). Tertiary alcohols ($R^4 = -C(CH_3)_2OH$) can be obtained from VI and MeMgBr. These tertiary alcohols can also be deoxygenated using the method of Lau. Similarly, the thiophene IX can be prepared from ketone VIII.

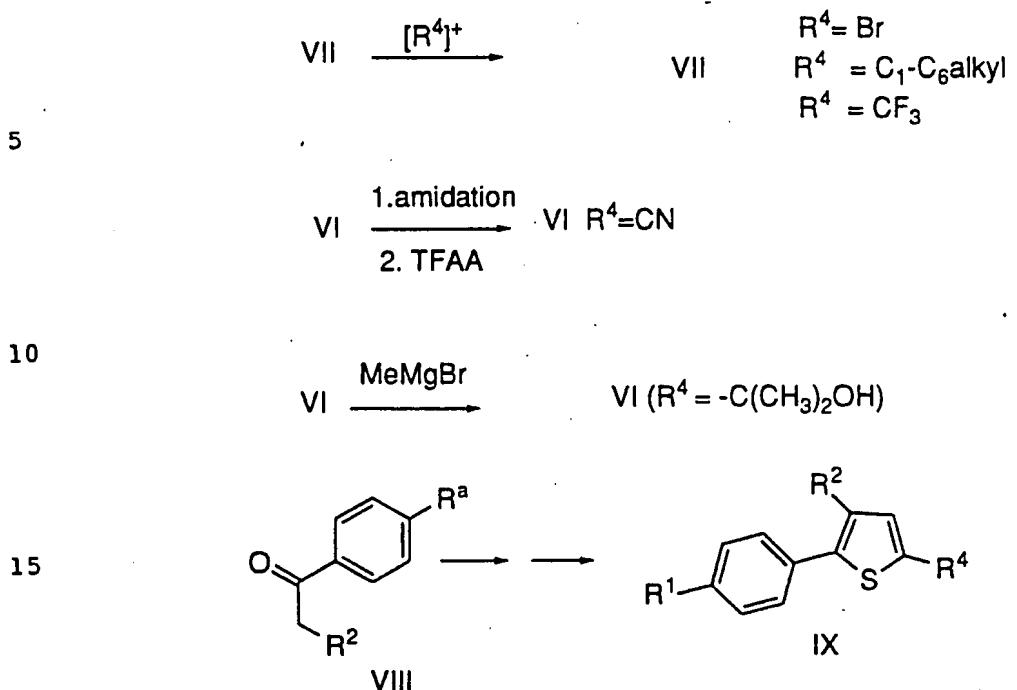
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METHOD A



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METHOD A CONT'D



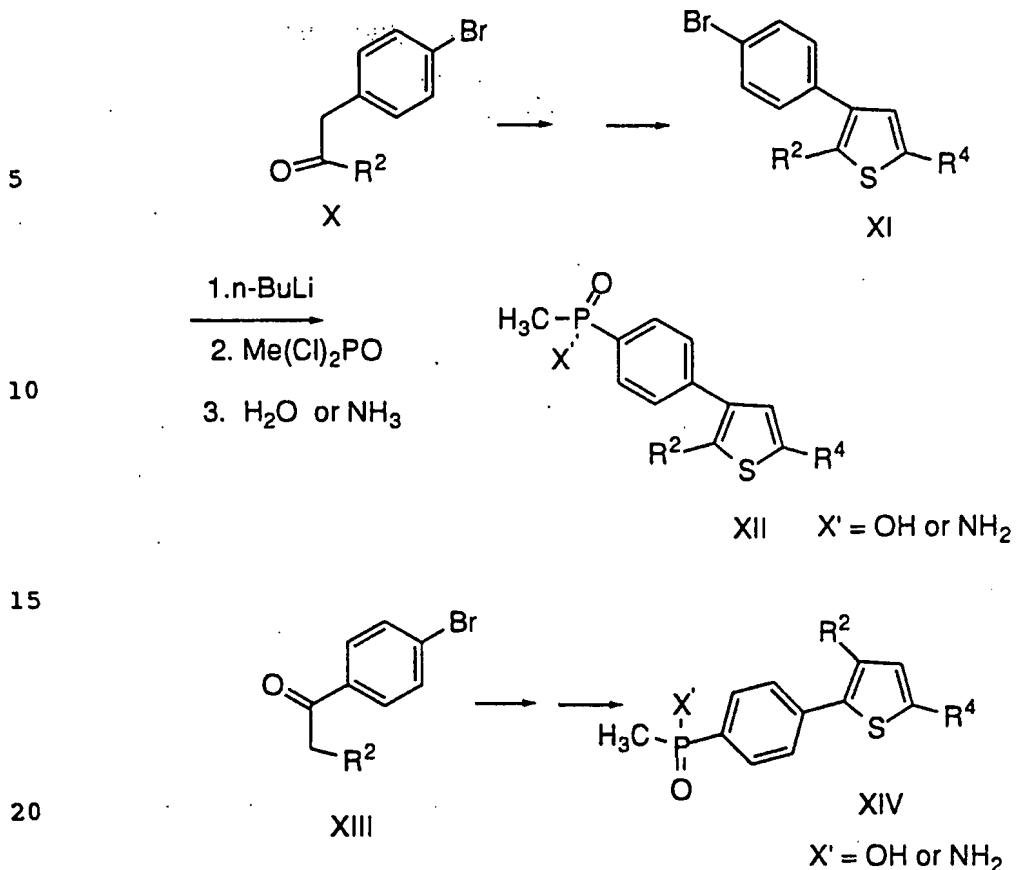
20 Method B: Ketone X can be converted to the thiophene compound XI using general methods already described in Method A. The thiophene XII can be prepared by metallation of XI with n-BuLi, quenching with methyl phosphonic dichloride and addition of water or ammonia ($X' = \text{OH}$ or NH_2). Similarly, the other regioisomer XIV can be prepared from ketone XIII.

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METHOD B



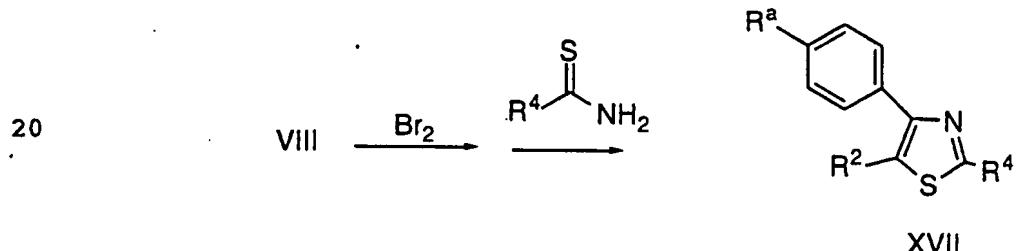
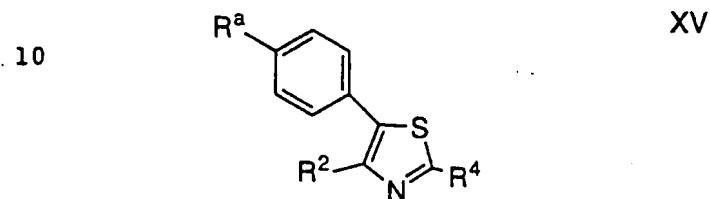
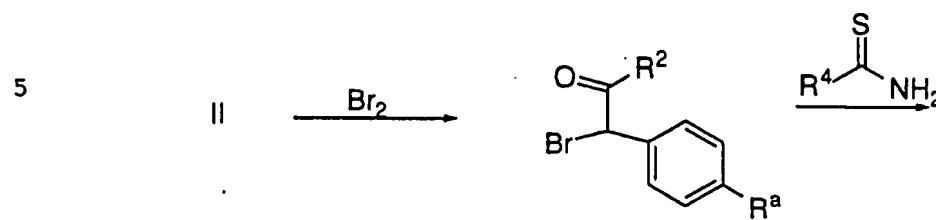
25 Method C:

Bromination of ketone II gives the α -bromoketone XV which is then converted to the thiazole XVI after treatment with a thioamide. Similarly, ketone VIII can be converted to thiazole XVII.

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METHOD C



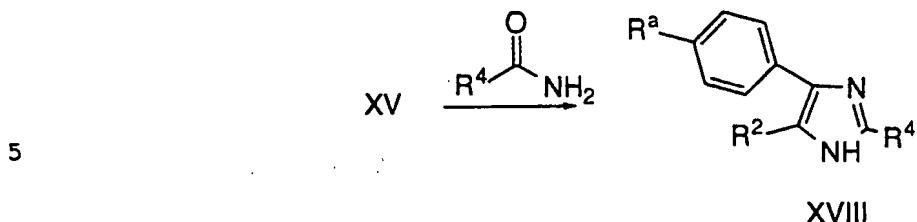
Method D:

25 Ketone XV can be converted to the imidazole compound XVIII after treatment with formamide using the preparation of Brederick et al, Chem. Ber. 1953, p. 88.

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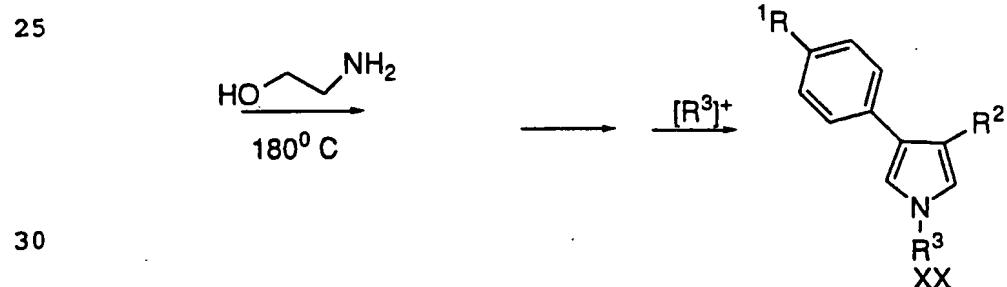
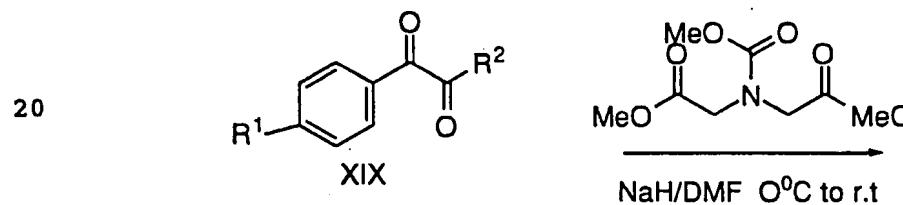
METHOD D

Method E:

10 Pyrole compound XX can be obtained from diketone XIX using the general procedures of Friedman et al, J. Org. Chem. 1965, 30, p. 854, K. Dimroth et al, Ber. 1956, 56, 2602, K. Dimroth et al, Ann. 1961, 634, 102. The free NH of the pyrole can be acylated with Cl-CO-lower alkyl in the presence of a base such as Et₃N. Also alkylated products can be prepared using alkyl halides as reagents with a base such as NaH.

15

METHOD E



Method F:

The compounds of type XXV can be prepared from readily available 4-substituted phenylacetyl chlorides XXIa. Reaction of di(3-but enyl)cadmium with a 4-substituted phenylacetyl chloride provides ketone XXI. Ozonolysis of XXI affords keto aldehyde XXIb which is cyclized by base to give cyclopentenone XXII. Addition of arylmagnesium bromide or aryllithium to XXII gives allylic alcohol XXIV. Oxidation of XXIV with pyridinium chlorochromate affords the desired 2,3-disubstituted cyclopentenone XXV. For preparation of compound XXV (R¹=SO₂Me), 4-methylthiophenyllithium is used followed by oxidation with the magnesium salt of monoperoxyphthalic acid (MMPP) or m-chloroperoxybenzoic acid (mCPBA) to introduce the required methylsulfonyl group in XXV.

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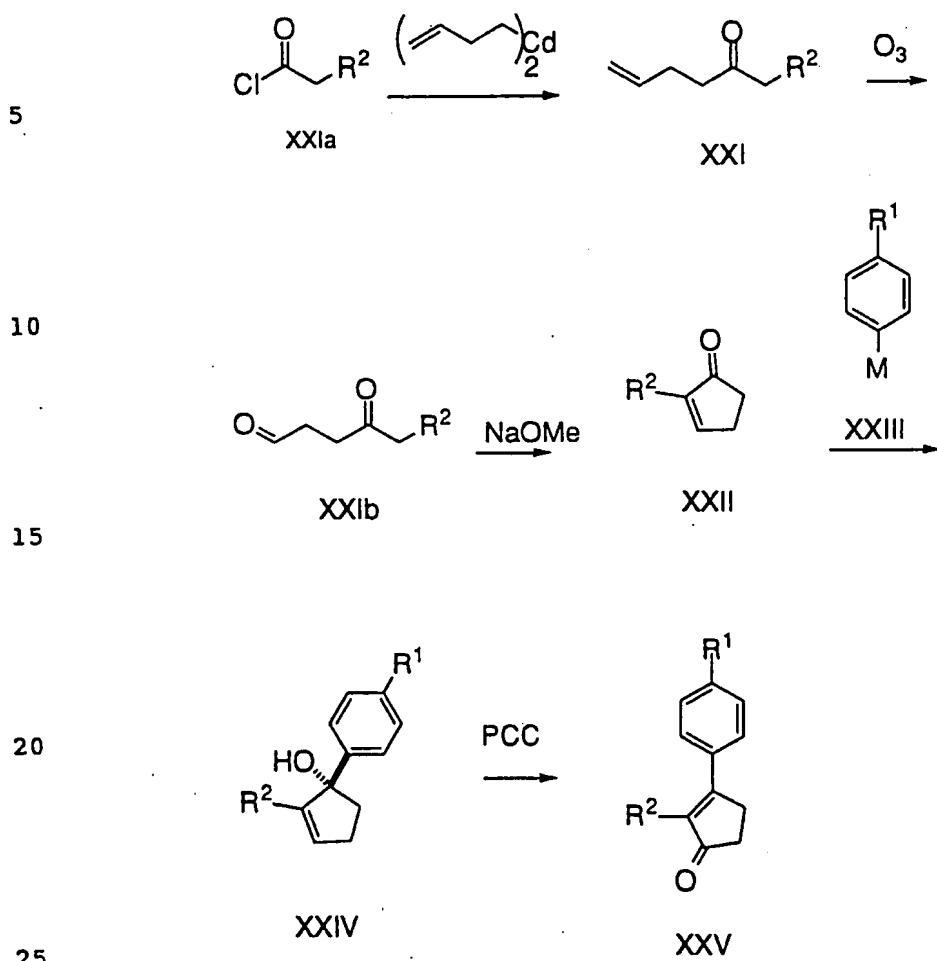
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METHOD F

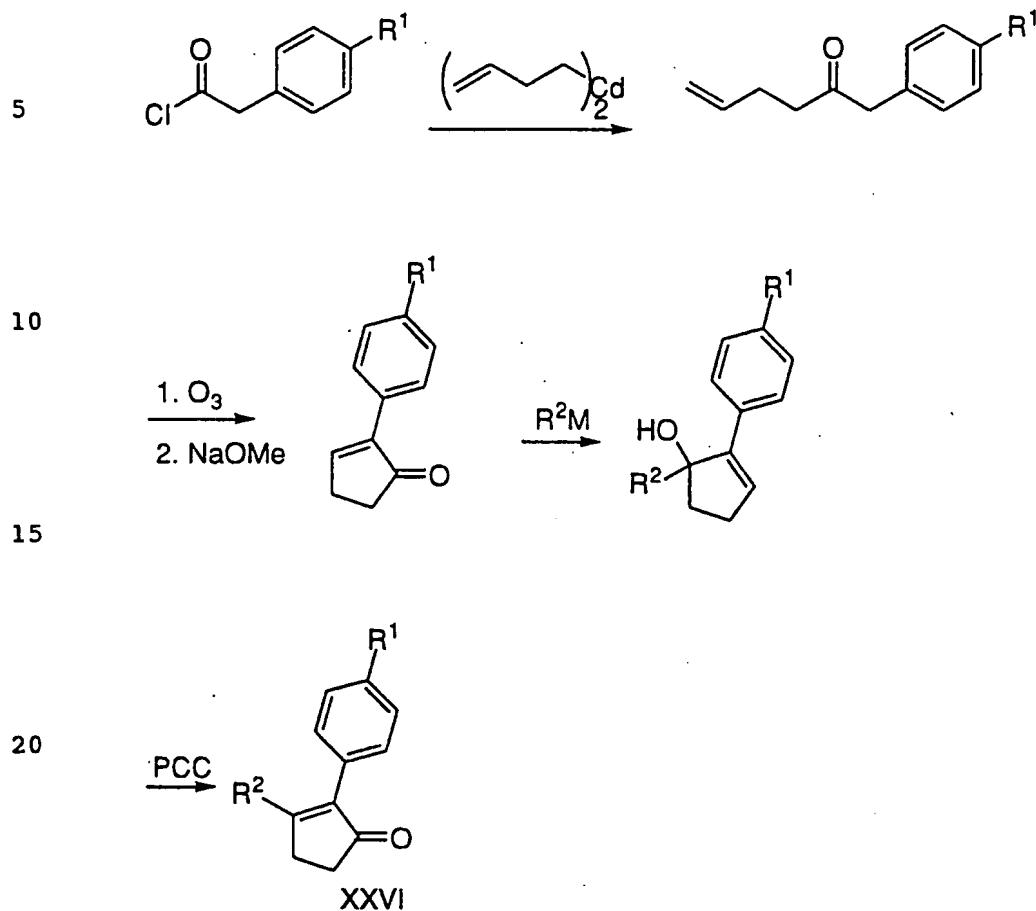
Method G:

The sequence of Method G is the same as in Method F except R¹ containing acid chloride is used as starting material. R² is introduced at a later stage via a carbonyl addition reaction, followed by PCC oxidation.

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METHOD G

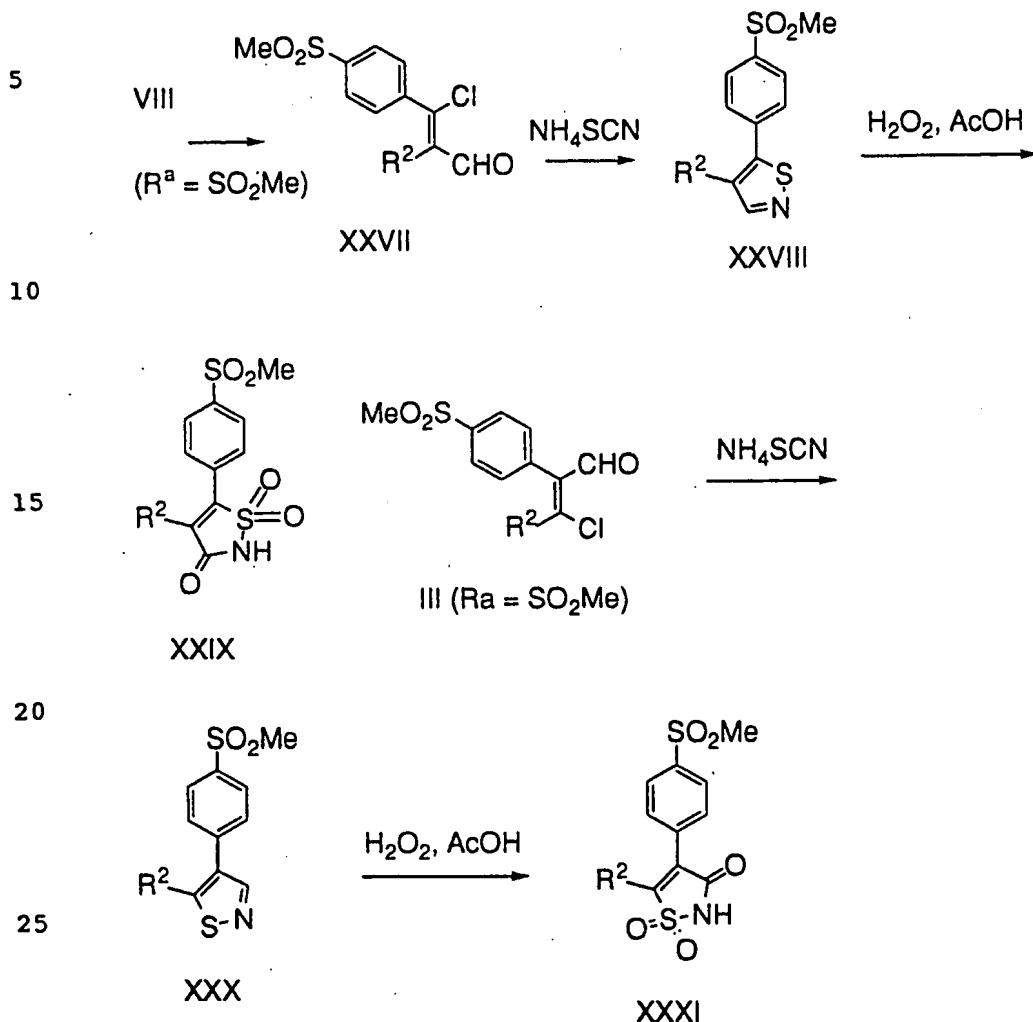
25 Method H:

The 4,5-disubstituted isothiazoles and isothiazol-3(2H)-one-1,1-dioxides can be prepared by the general method described by B. Schulze et al, Helvetica Chimica Acta, 1991, 74, 1059. Thus, aldehyde III ($R^a=SO_2Me$) or XXVII is treated with excess NH_4SCN in refluxing acetone to provide the corresponding 4,5-disubstituted isothiazoles XXX

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and XXVIII, oxidation of which with hydrogen peroxide yields XXXI and XXIX.

METHOD H



Method I:

30 An appropriately substituted aryl bromomethyl ketone is reacted with an appropriately substituted aryl acetic acid in a solvent such

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as acetonitrile in the presence of a base such as triethylamine and then treated with 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) to afford either the lactone XXXIII or XXXV.

METHOD I

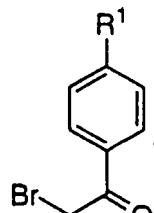
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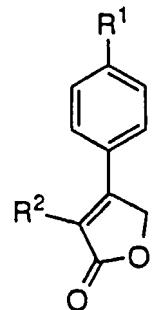
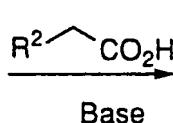
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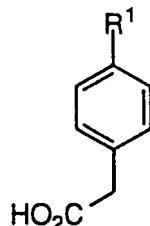
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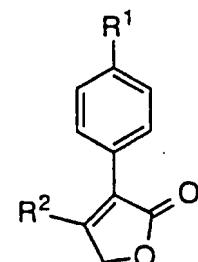
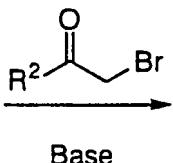
XXXII



XXXIII



XXXIV



XXXV

(R² is a mono- or disubstituted phenyl or
a mono- or disubstituted heteroaryl)

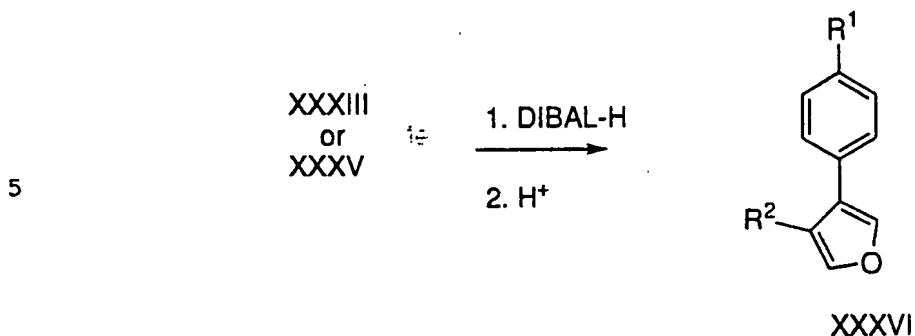
Method J:

Either of the lactones XXXIII or XXXV in a solvent such as THF is reacted with a reducing agent such as diisobutyl aluminium hydride or lithium borohydride at -78°C, to yield the furan XXXVI.

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METHOD J



10 Method K:

The preparation of lactams XXXVII and XXXIX can be achieved by the same reaction as described in Method I, except an appropriate amide XXXVIII is used.

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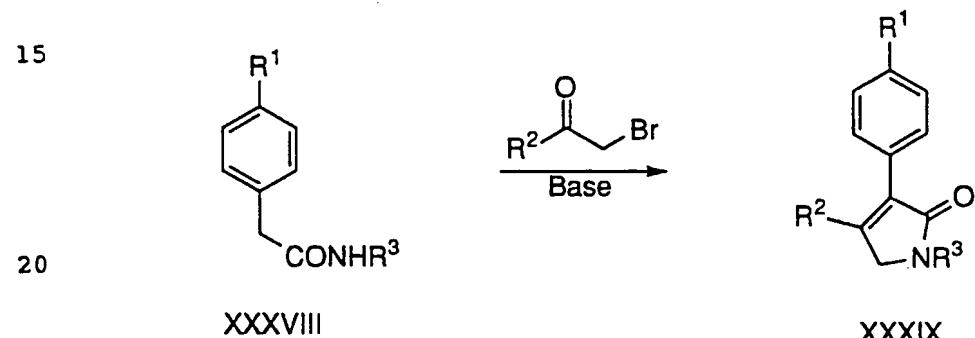
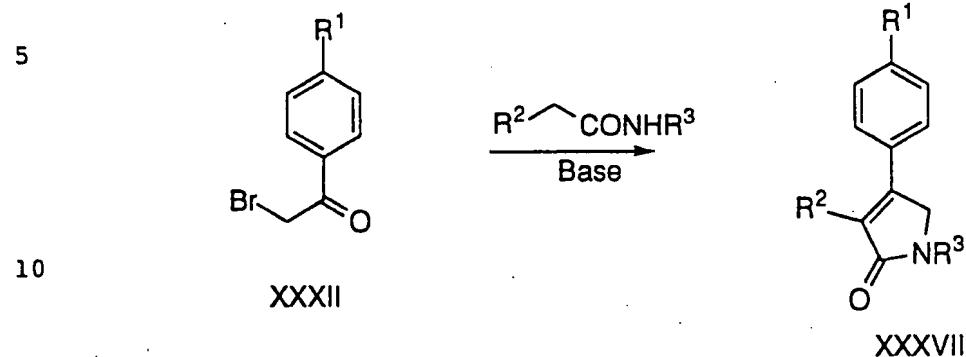
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METHOD K

Method L:

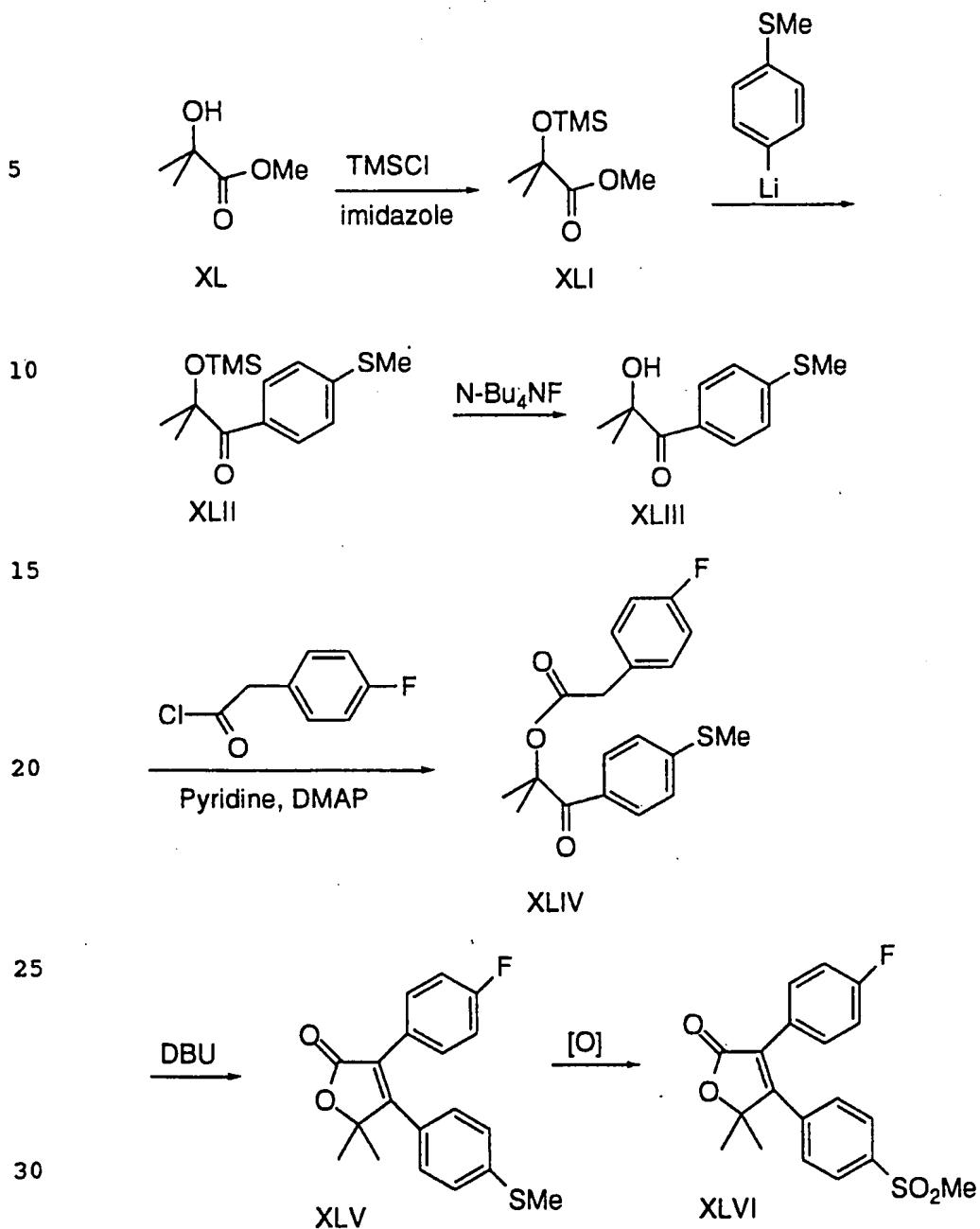
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Methyl 2-hydroxy isobutyrate is silylated with TMSCl to give the TMS ether XLI, which is treated with 4-methylthiophenyllithium to provide ketone XLII. Desilylation followed by acylation yields keto-ester XLIV, which can be cyclized to lactone XLV by base catalysis. Oxidation of XLV with MMPP or mCPBA affords the desired product XLVI.

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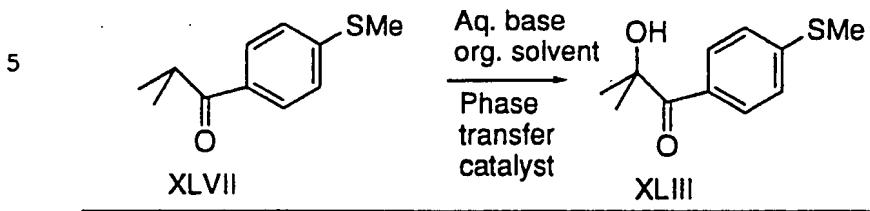
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METHOD L



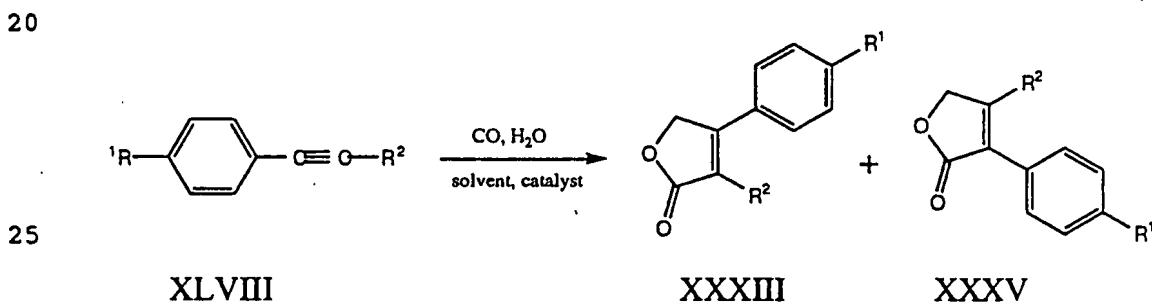
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METHOD M



10 An alternative preparation of the hydroxy ketone XLIII is the oxidation of the known (J. Org. Chem. 1991 **56**, 5955-8; Sulfur Lett. 1991, **12**, 123-32) ketone XLVII. A mixture of XLVII, aqueous base, such as NaOH, organic solvents such as carbon tetrachloride/toluene and a phase transfer catalyst such as ALIQUAT 336 is stirred in air at room
 15 temperature to provide XLIII. Compound XLIII is also described in U.S. 4,321,118 and Org. Coat. 1986, **6**, 175-95.

Method N

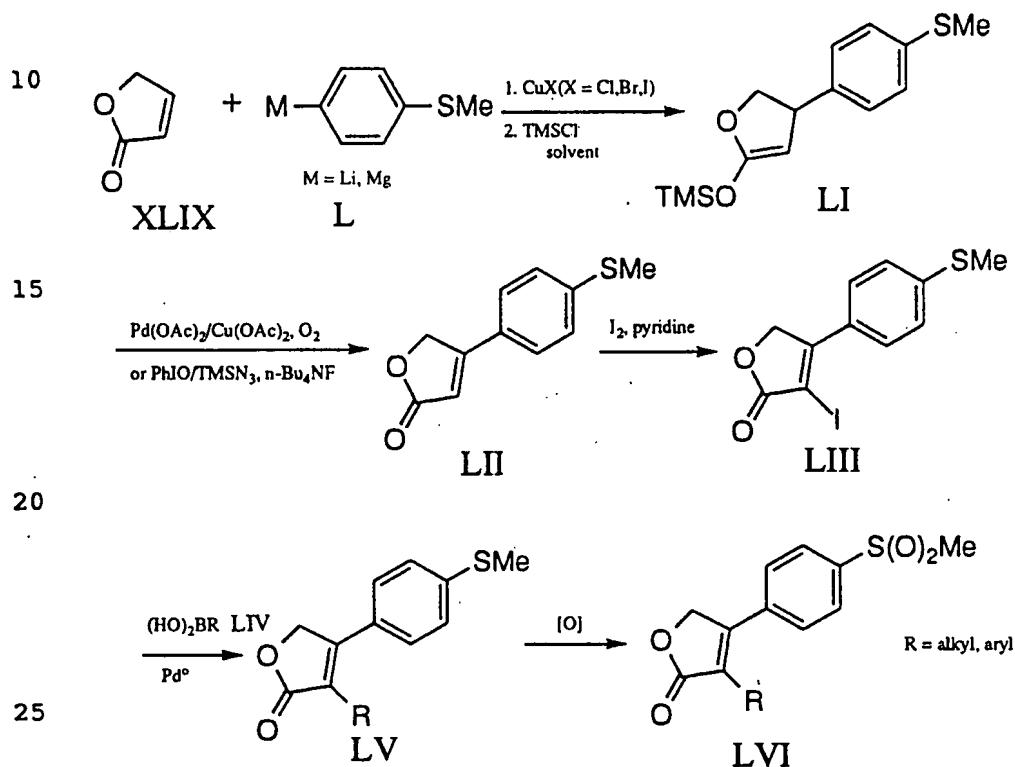


30 By reacting an acetylene XLVIII with carbon monoxide and water in the presence of suitable catalysts, a mixture of compound XXXIII and its isomer XXXV is obtained. The isomers are separable by standard procedures in the art such as chromatography or crystallization. Examples of useful catalysts and conditions are PdCl_2 in aqueous HCl and EtOH ,

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heated at 50-150°C and 50-150 atmospheres of pressure, or Rh₄(CO)₁₂ (or Rh₆(CO)₁₆) in aqueous THF (or acetone, acetonitrile, benzene, toluene, EtOH, MeOH) containing a trialkylamine, at 50-150°C and 20-300 atmospheres pressure. See Takahashi et al., *Organometallics* 1991, 10, 2493-2498; and Tsuji et. al., *J. Am. Chem. Soc.* 1966, 88, 1289-1292.

Method O



1, 4-Addition to XLIX of 4-methylthiophenyl organometallic reagents L in the presence of copper salts and the trapping of the resultant enolate with trialkyl silyl chloride such as TMSCl or TIPSCl provide the ketene acetal LI. The ketene acetal can then be oxidized to the substituted butenolide LII by the method of Ito using catalytic amounts of Pd₂(OAc)₂ and Cu(OAc)₂ and O₂ in MeOH or by the method of Magnus using

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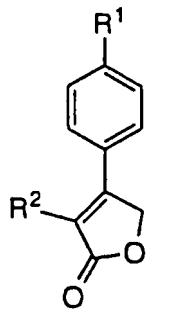
and Cu(OAc)₂ and O₂ in MeOH or by the method of Magnus using PhIO/TMSN₃ and Bu₄NF. Introduction of the iodine can be accomplished by treating LII with I₂ in the presence of pyridine to afford LIII.

5 Palladium catalyzed Susuki or Stille coupling of LIII with the appropriate aryl or alkyl partner such as the boronic acid LIV provides the butenolide LV. The sulfide can be oxidized to a sulfone by various oxidizing agents such as peracetic acid, MPPM, MMPP or H₂O₂ to give the desired compound LVI. See Y. Ito et. al., *J. Am. Chem. Soc.* 1979, 101, 494; and P. Magnus et. al., *Tet. Lett.* 1992, 2933.

10 Accordingly, in a further aspect the invention is directed to a process of making a compound of formula XXXIII

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comprising:

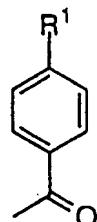
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(a1) reacting in an organic solvent a compound of formula XXXII'

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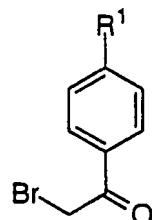


XXXII'

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with a bromine reagent to yield a compound of formula XXXII

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XXXII

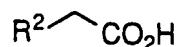
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For purposes of this specification the organic solvent shall be defined to include, but not be limited to methylene chloride, chloroform, carbontetrachloride and acetic acid. Similarly, the bromine reagent shall be defined to include, but not be limited to bromine, pyridinium perbromide hydrobromide, CuBr₂, and N-bromosuccinimide.

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(a2) reacting in a non-aqueous polar solvent a compound of formula XXXII

with a compound of formula

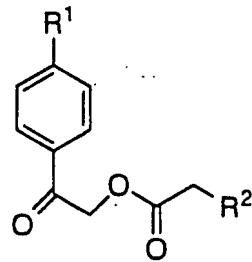


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in the presence of a base to produce a compound of formula A

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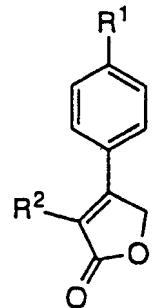
A

(a3) treating in a non-aqueous polar solvent a compound of formula A with
10 strong base to yield a compound of formula XXXIII.

15 For purposes of this specification the non-aqueous polar solvent shall be defined to include, but not be limited to, acetonitrile propionitrile, acetone, 2-butanone and tetrahydrofuran. Similarly, the base is defined to include, but not be limited to a tri-C₁-3alkylamine such as triethylamine. Moreover, the strong base is defined to include, but not be limited to, an amidine, a guanidine, lithium diisopropylamide and potassium bis-(trimethylsilyl) amide.

20 In an alternative, the invention is directed to a process of making a compound of formula XXXIII

25



30

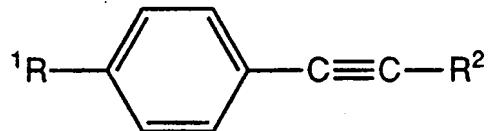
XXXIII

- 52 -

comprising:

(b1) reacting an acetylene compound of the formula XLVIII

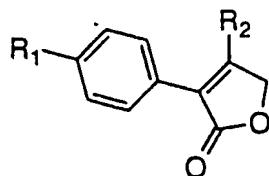
5



XLVIII

10 with carbon monoxide and water in the presence of a suitable catalyst to
10 yield a compound of formula XXXIII and XXXV.

15



XXXV

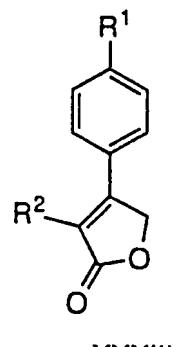
20 For purposes of this specification suitable catalysts include, but
20 are not limited to $\text{Ru}_4(\text{CO})_{12}$, $\text{Co}_2(\text{CO})_8$ or PdCl_2 in aqueous THF or
acetone, acetonitrile, benzene, toluene, methyl alcohol or ethyl alcohol.

25 In a second alternative, the invention is directed to a process of
25 making a compound of formula XXXIII

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- 53 -

5

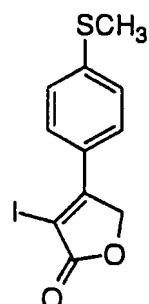


10 comprising:

(c1) reacting a compound of formula LIII

15

20

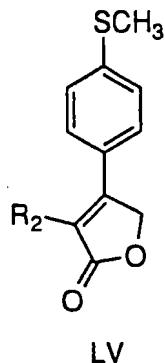


25 with a reagent of the formula $(HO)_2BR^2$ in an aqueous solvent such as benzene, toluene, THF, MeOH, DME or EtOH and in the presence of a suitable palladium catalyst to yield a compound of formula LV, and

30

- 54 -

5



LV

10 (c2) oxidizing the compound of formula LV to yield a compound of formula XXXIII.

15 For purposes of this specification, the catalyst is defined to include, but not be limited to palladium catalysts. Similarly, the solvent is intended to include, but not be limited to benzene, toluene, THF, MeOH, DME or EtOH.

20 In all of the process alternatives, R1 and R2 are as defined above for the portion of Detailed Description and Claims directed to the compounds of formula I.

Representative Compounds

Tables I and II illustrate compounds of formula I.

25

30

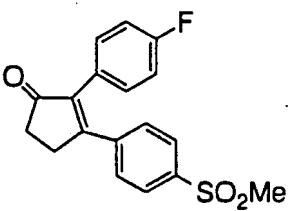
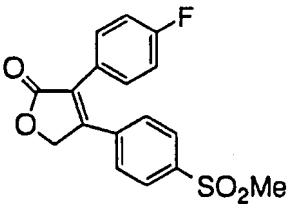
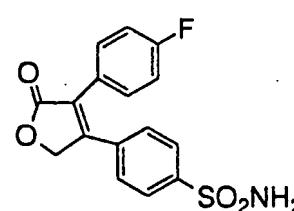
- 55 -

Table I

		Example	Method
5		1	A
10		2	A
15		3	A
20		4	A
25		5	A
30		6	C

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Table I (continued)

		Example	Method
5		7	F
10		8	H
15		9	I
20		10	I
25			
30			

- 57 -

Table I (continued)

Example Method

5		11	J
10		12	L
15		13	A
20		14	A
25			
30			

- 58 -

Table I (continued)

		Example	Method
5		15	I
10		16	I
15		17	I
20		18	I
25		19	I
30			

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Table I (continued)

		Example	Method
5		20	1
10		21	1
15		22	1
20		23	1
25		24	1
30			

- 60 -

Table I (continued)

		Example	Method
5		25	
10		26	
15		27	
20		28	
25		29	
30			

- 61 -

Table I (continued)

		Example	Method
5		30	1
10		31	1
15		32	1
20		33	1
25		34	1
30			

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Table I (continued)

		Example	Method
5		35	
10		36	
15		37	
20		38	
25		39	
30			

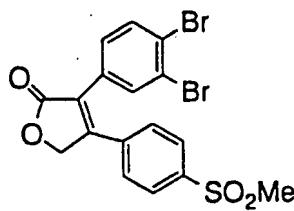
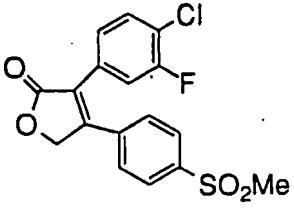
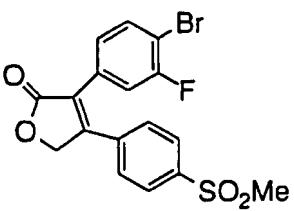
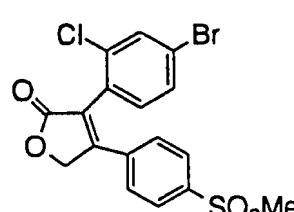
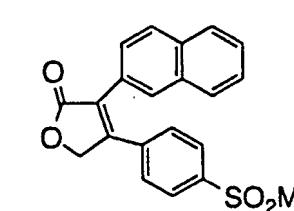
- 63 -

Table I (continued)

		Example	Method
5		40	I
10		41	I
15		42	I
20		43	I
25		44	I
30			

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Table I (continued)

		Example	Method
5		45	
10		46	
15		47	
20		48	
25		49	
30			

- 65 -

Table I (continued)

		Example	Method
5		50	I
10		51	I
15		52	I
20		53	I
25		54	I
30			

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Table I (continued)

Example Method

5		55	H
10		56	L + M
15		57	L + M
20		58	L + M
25			
30			

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Table I (continued)

Example Method

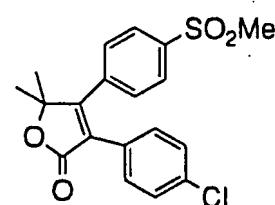
5



59

L + M

10



60

L + M

15

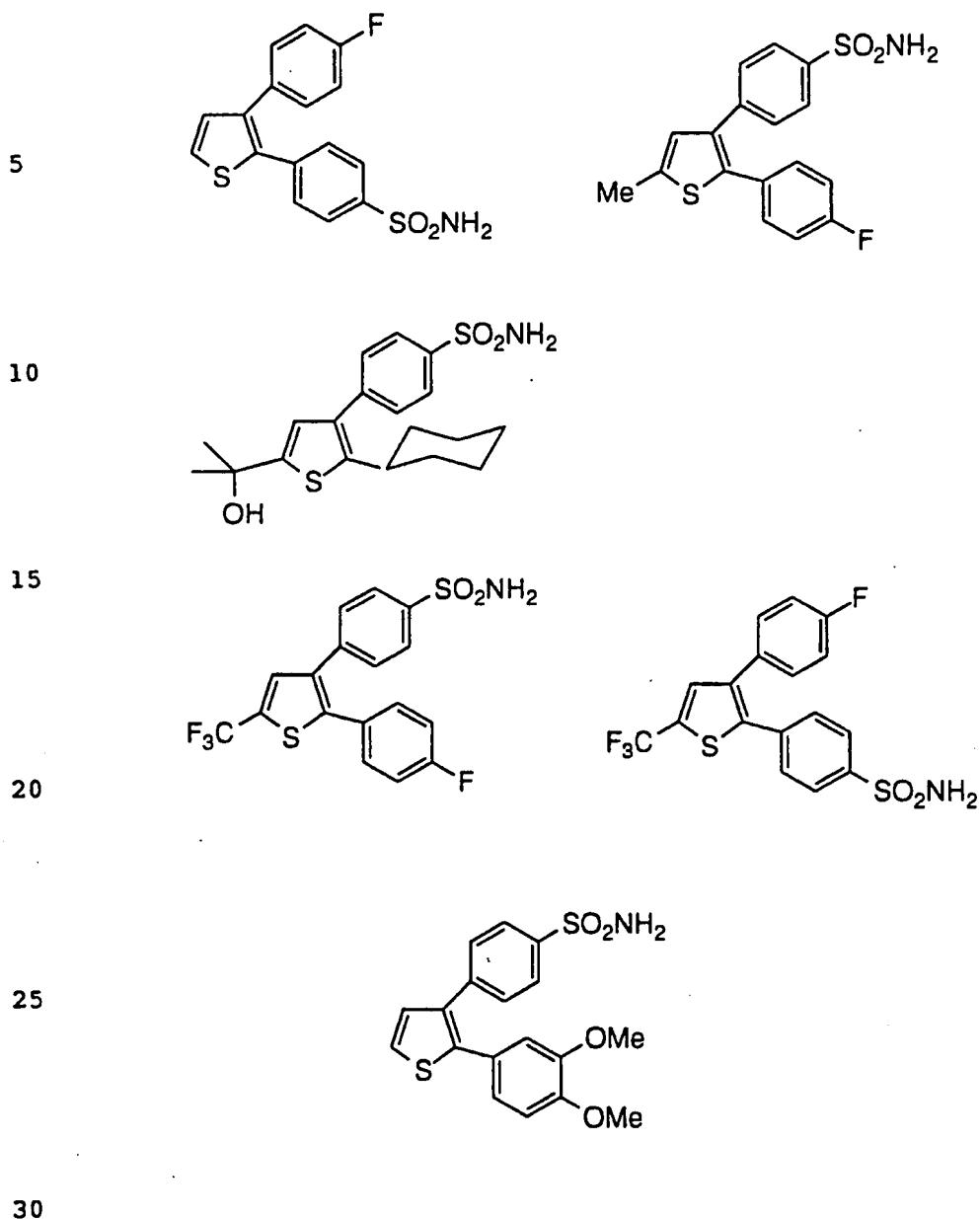
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25

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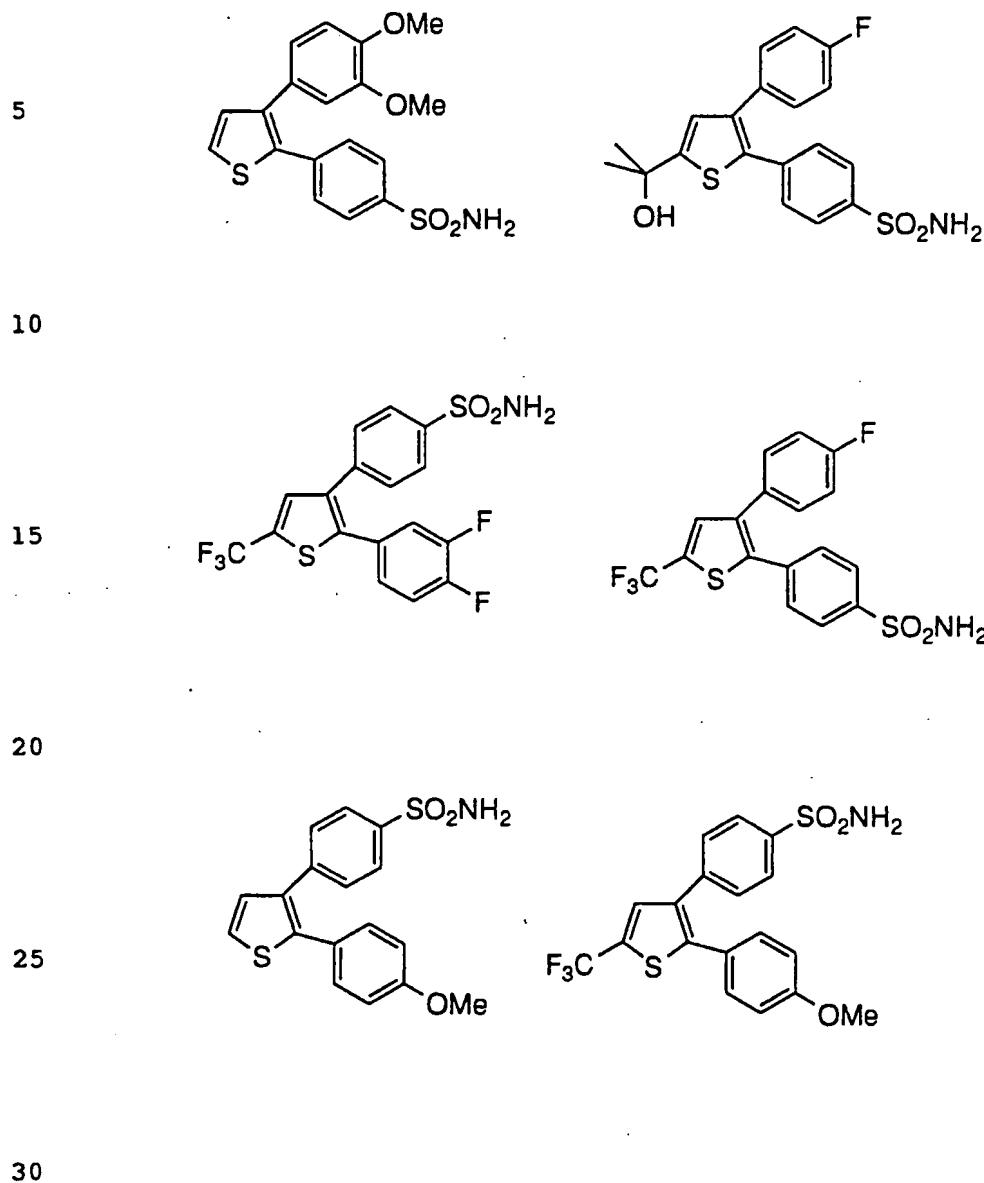
- 68 -

Table II



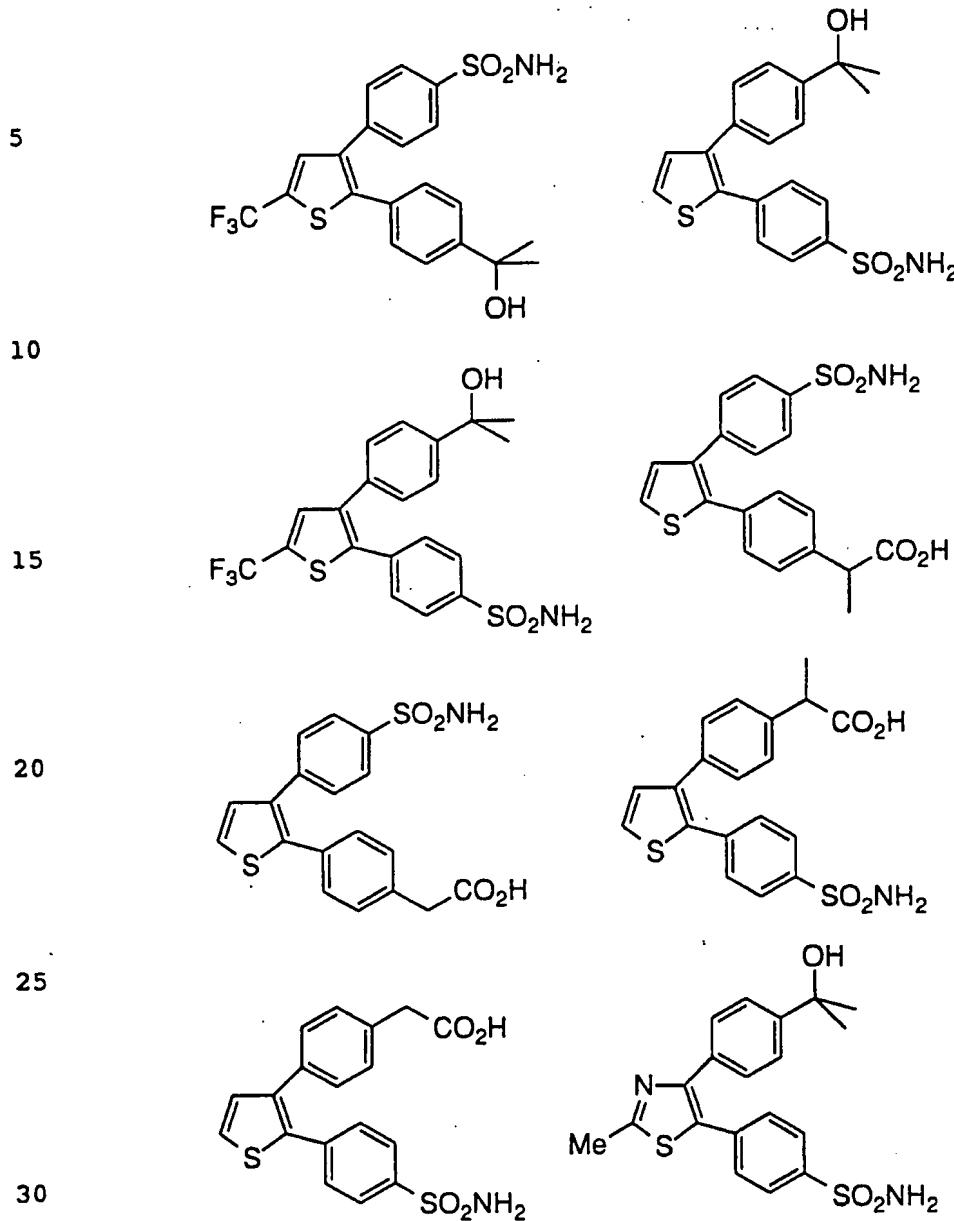
- 69 -

Table II (continued)



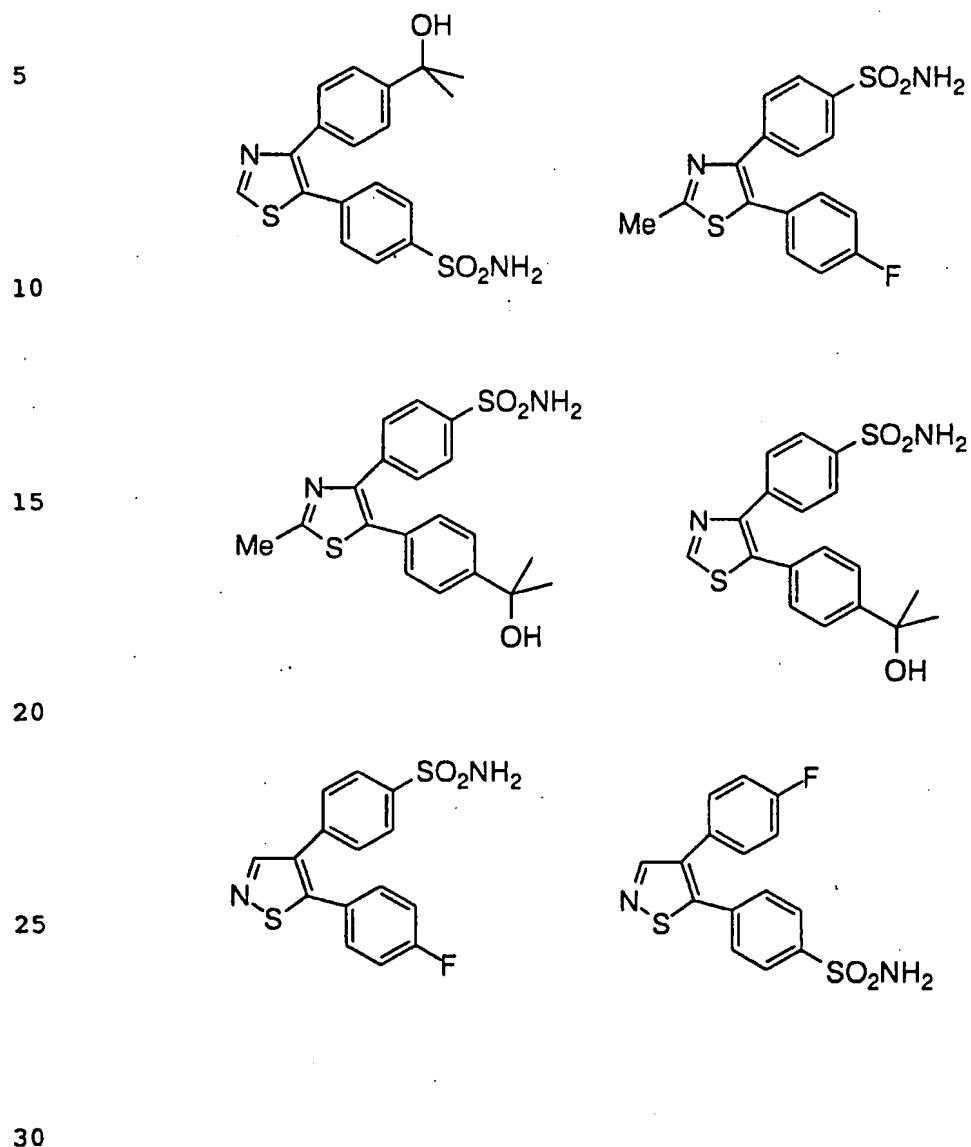
- 70 -

Table II (continued)



- 71 -

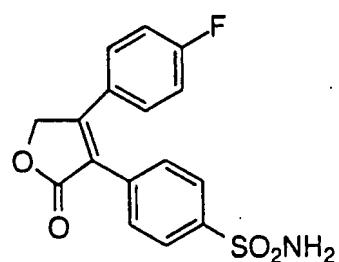
Table II (continued)



- 72 -

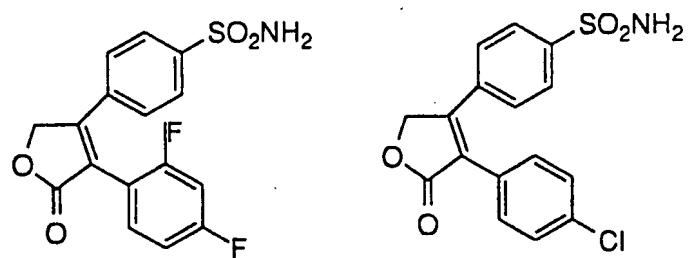
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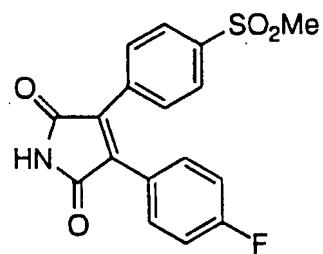
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15



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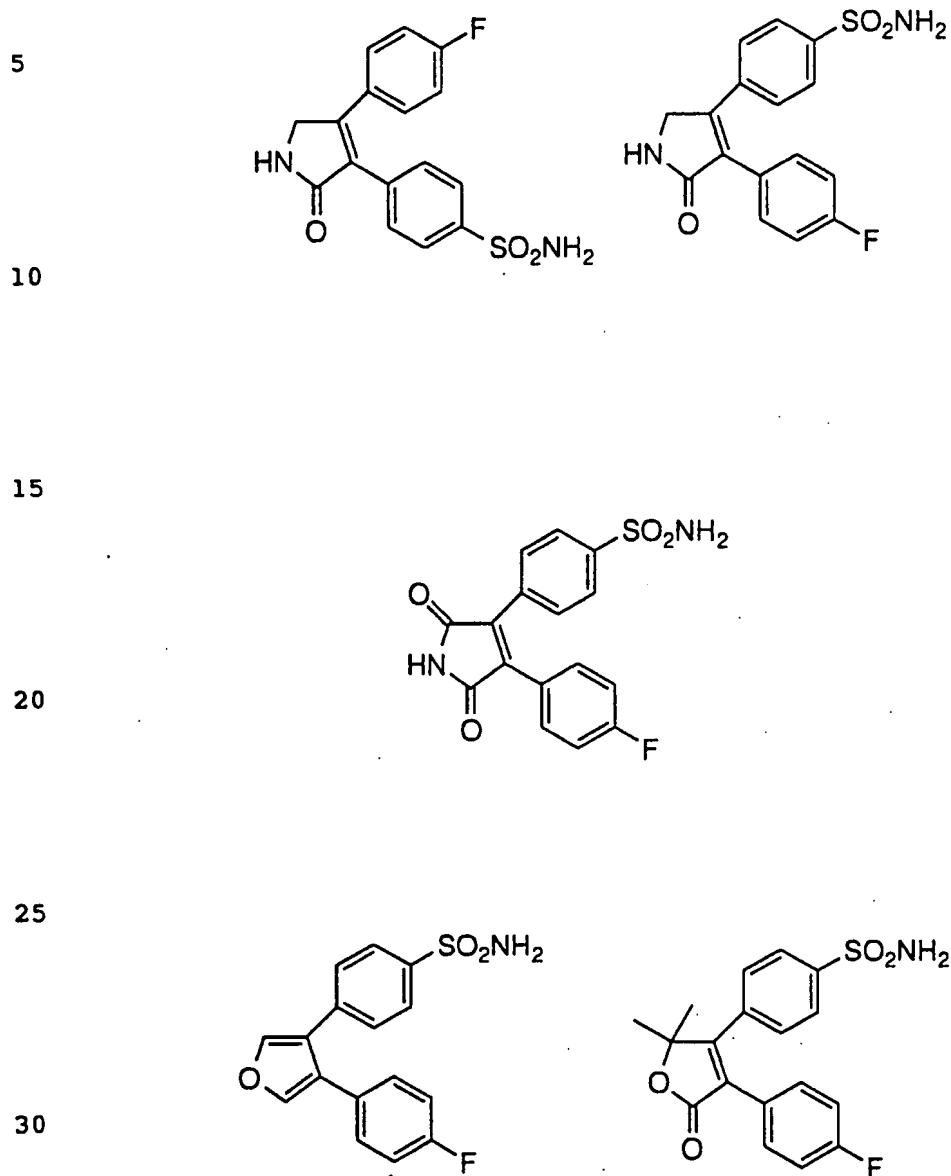
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30

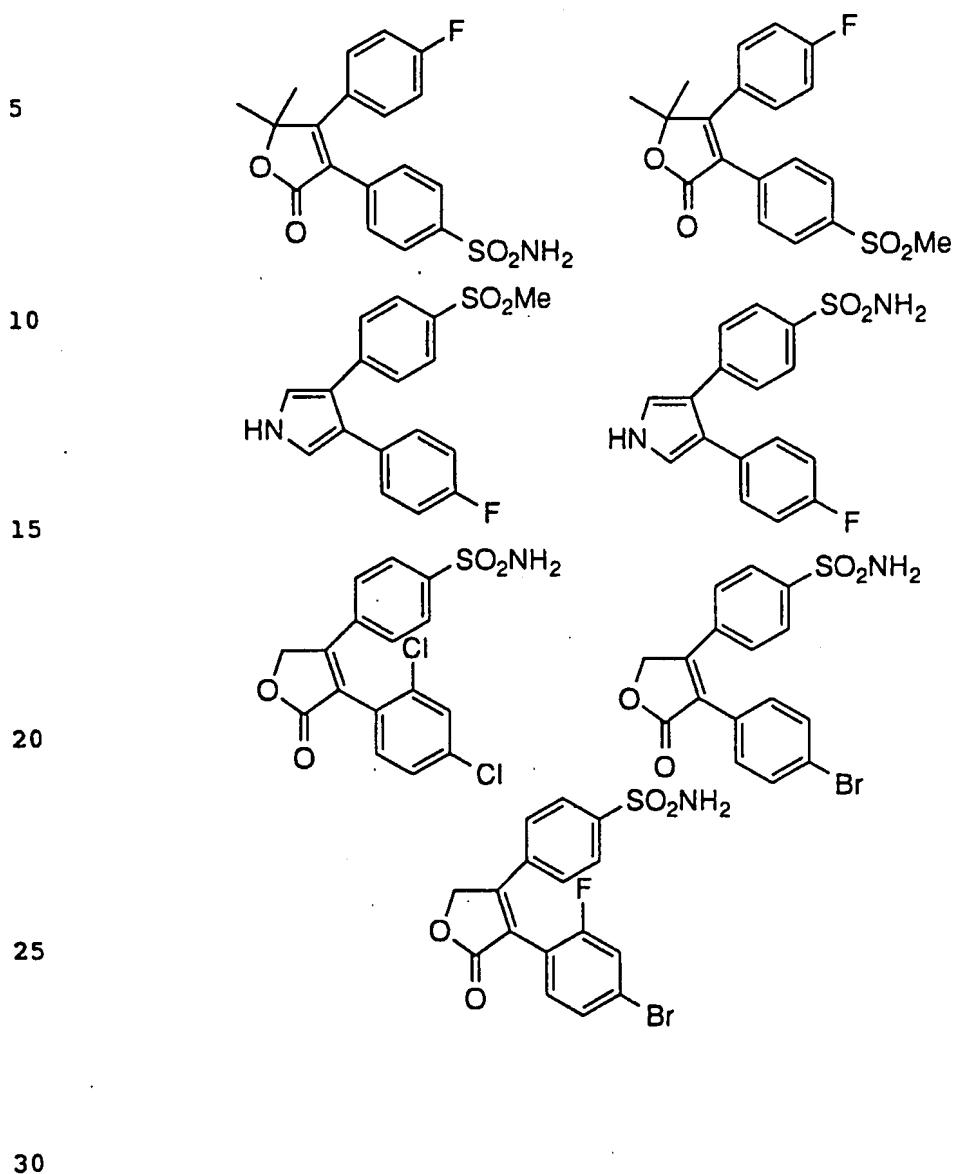
- 73 -

Table II (continued)



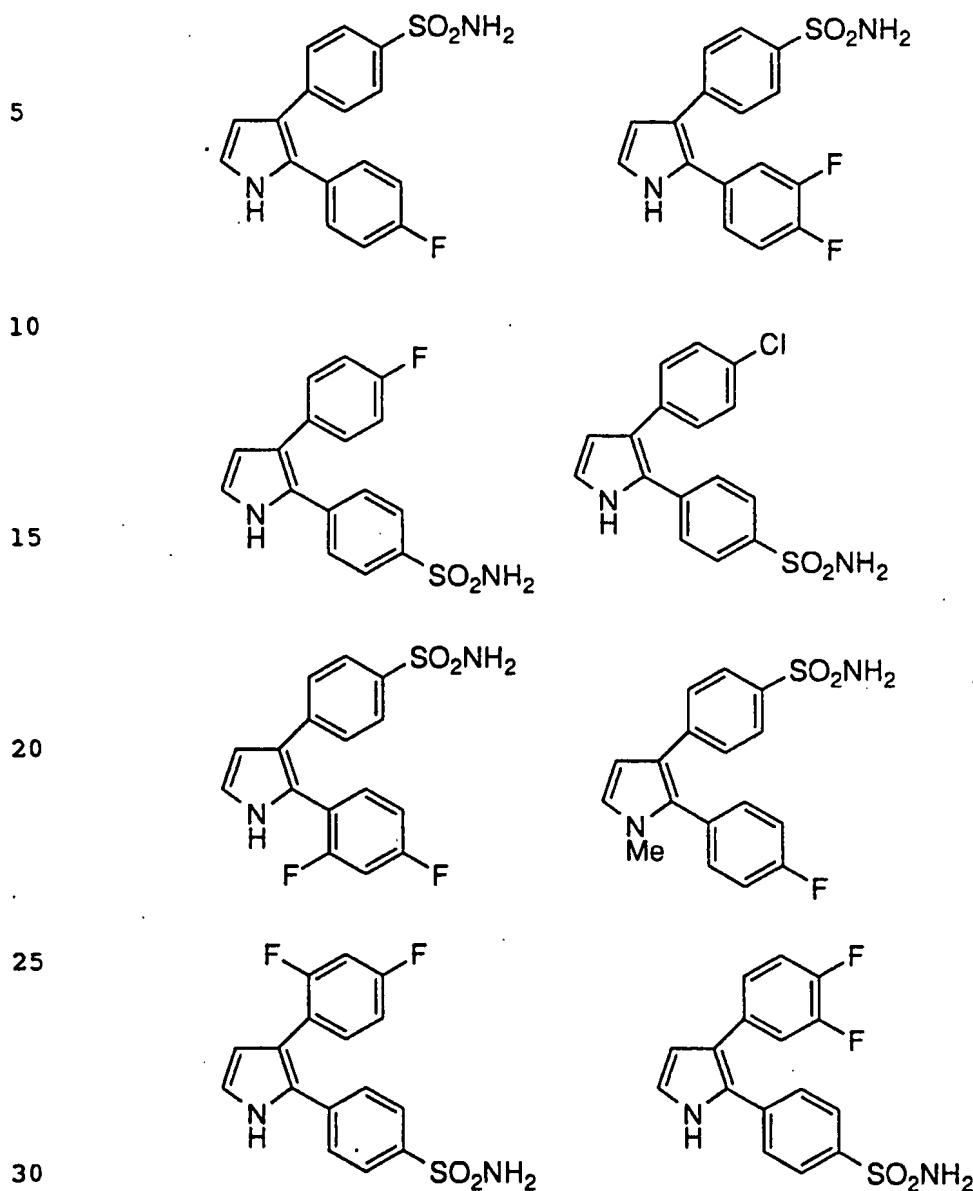
- 74 -

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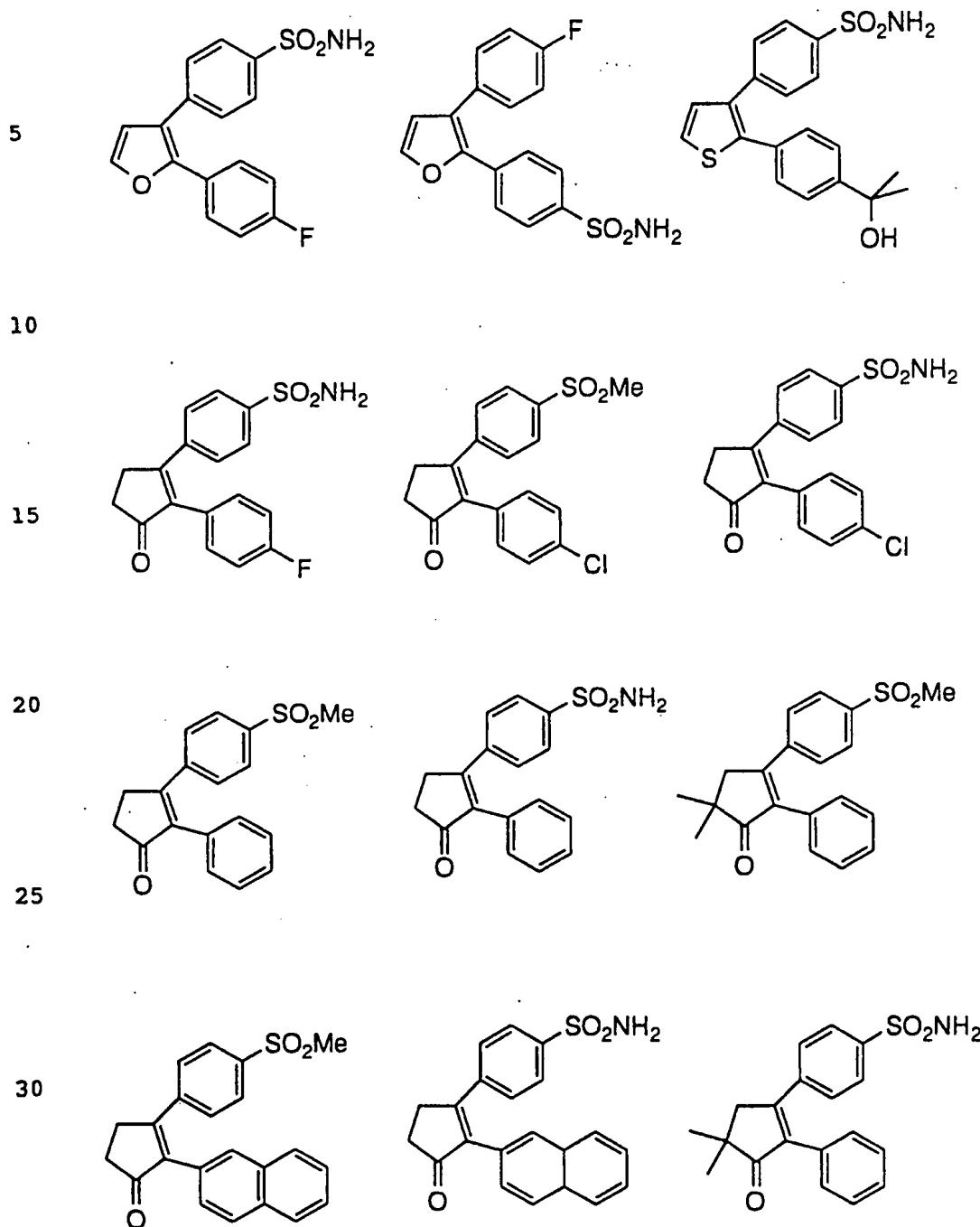
- 75 -

Table II (continued)



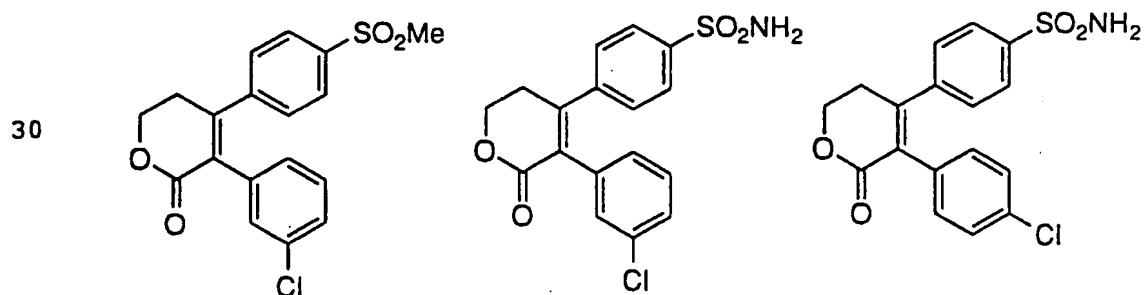
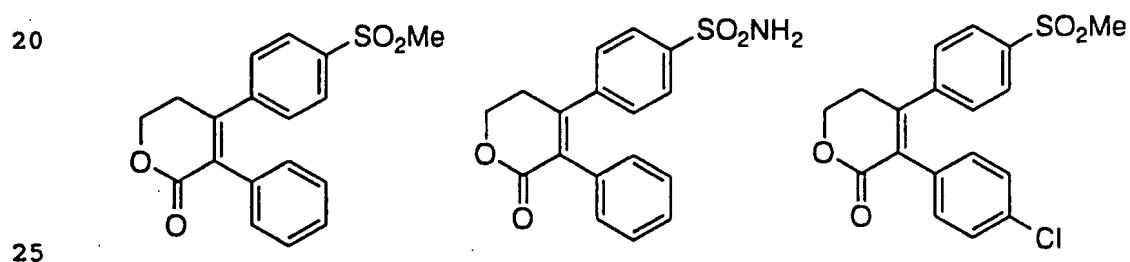
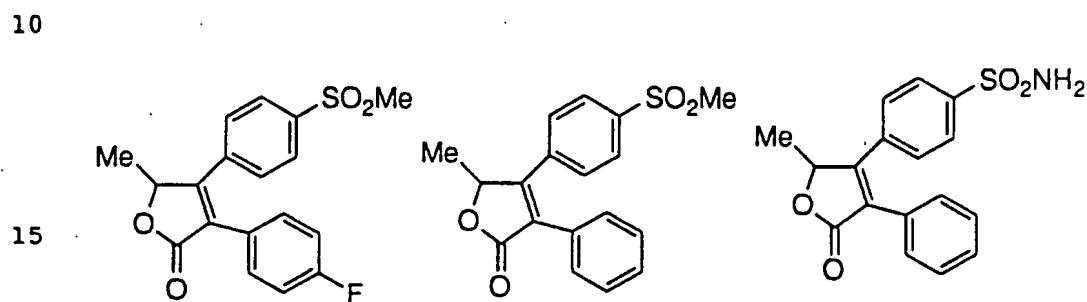
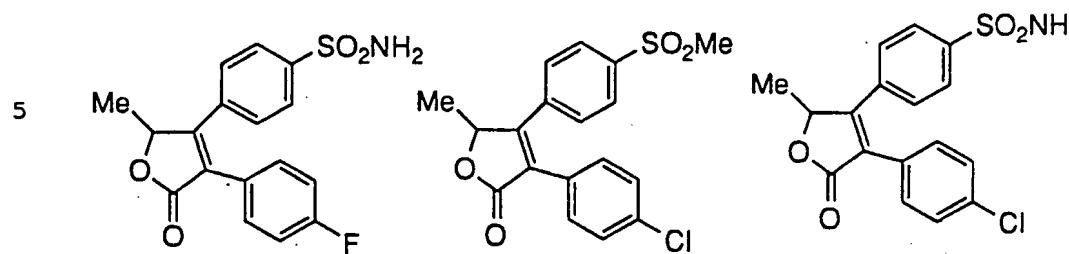
- 76 -

Table II (concluded)



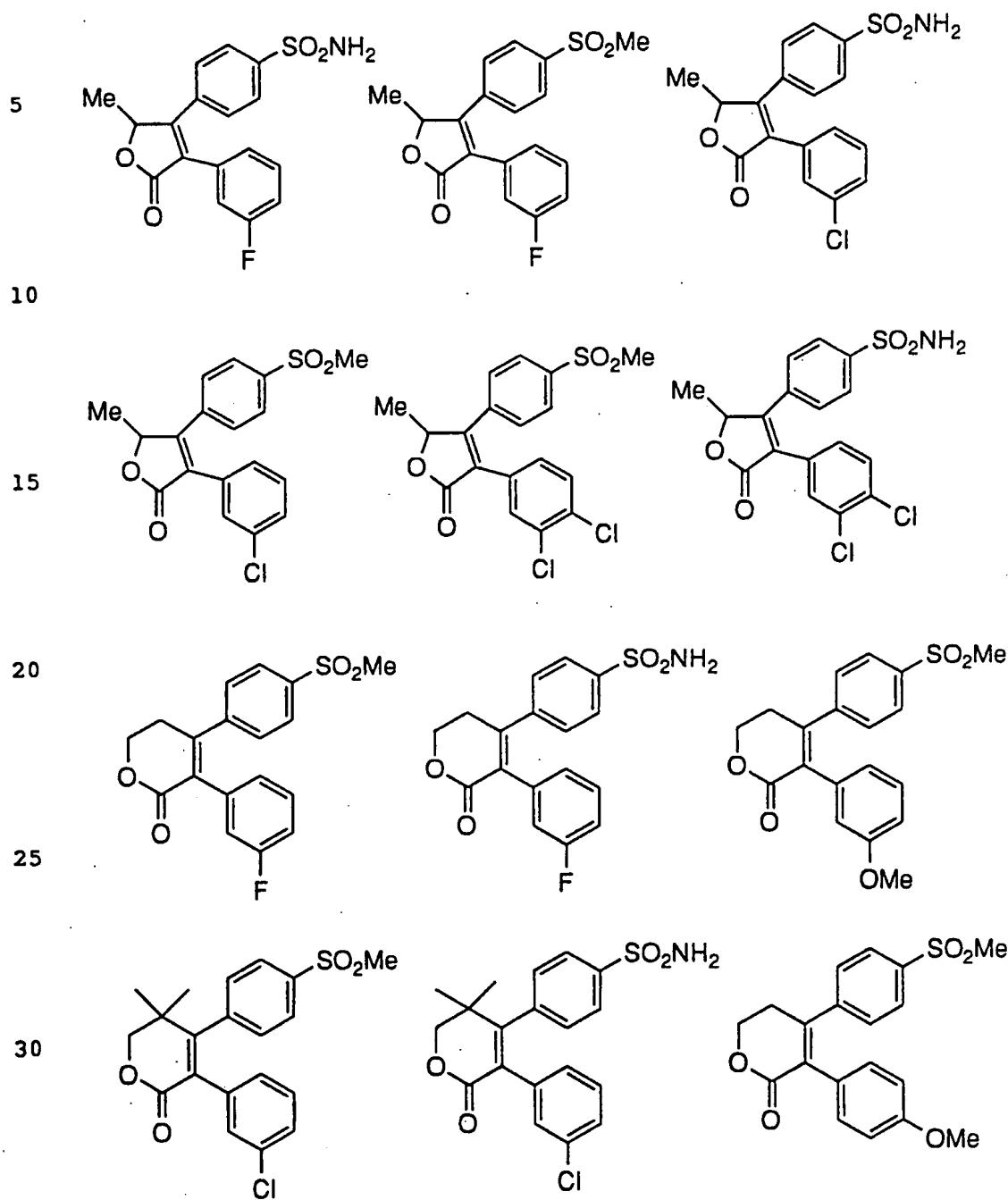
- 77 -

Table II (concluded)



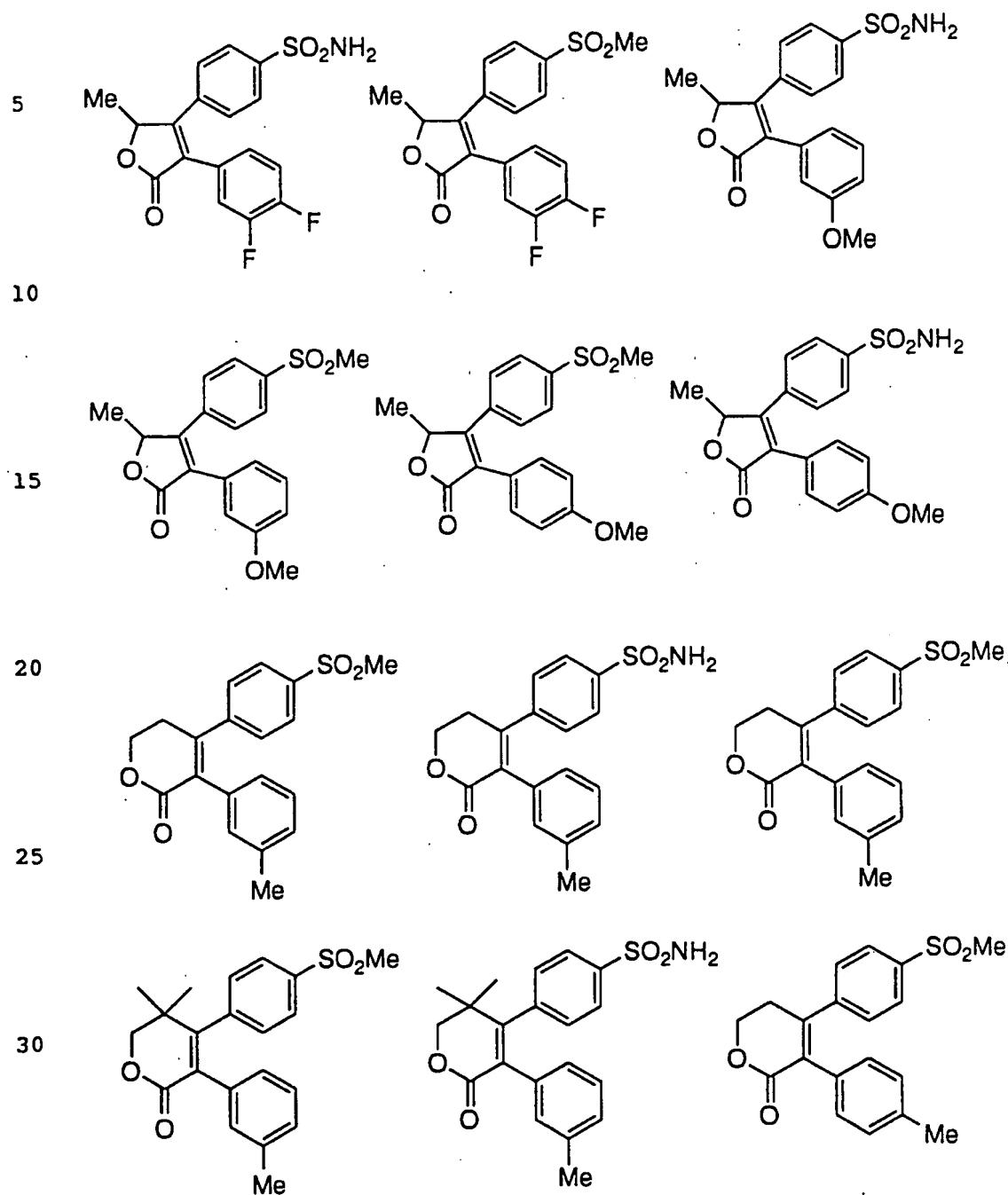
- 78 -

Table II (concluded)



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Table II (concluded)



Assays for Determining Biological Activity

The compound of Formula I can be tested using the following assays to determine their cyclooxygenase-2 inhibiting activity.

5 Inhibition of Cyclooxygenase Activity

Compounds were tested as inhibitors of cyclooxygenase activity in whole cell and microsomal cyclooxygenase assays. Both of these assays measured prostaglandin E2 (PGE2) synthesis in response to arachidonic acid, using a radioimmunoassay. Cells used for whole cell assays, and from which microsomes were prepared for microsomal assays, 10 were human osteosarcoma 143 cells (which specifically express cyclooxygenase-2) and human U-937 cells (which specifically express cyclooxygenase-1). In these assays, 100% activity is defined as the difference between prostaglandin E2 synthesis in the absence and presence 15 of arachidonate addition. IC₅₀ values represent the concentration of putative inhibitor required to return PGE2 synthesis to 50% of that obtained as compared to the uninhibited control. Representative results are shown in Table III.

20 Representative Rat Paw Edema Assay – Protocol

Male Sprague-Dawley rats (150-200g) were fasted overnight and were given po either vehicle (5% tween 80 or 1% methocel) or a test compound at 9 - 10 am. One hr later, a line was drawn using a permanent 25 marker at the level above the ankle in one hind paw to define the area of the paw to be monitored. The paw volume (V_{0h}) was measured using a plethysmometer (Ugo-Basile, Italy) based on the principle of water displacement. The animals were then injected subplantarly with 50 ul of a 1% carrageenan solution in saline (FMC Corp, Maine) into the paw using 30 an insulin syringe with a 25-gauge needle (i.e. 500 ug carrageenan per paw). Three hr later, the paw volume (V_{3h}) was measured and the

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increases in paw volume ($V_{3h} - V_{0h}$) were calculated. The animals were euthanized by CO₂ asphyxiation and the absence or presence of stomach lesions scored. Stomach scores were expressed as the sum of total lesions in mm. Paw edema data were compared with the vehicle-control group and percent inhibition calculated taking the values in the control group as 100%. Since a maximum of 60 - 70% inhibition (paw edema) was obtained with standard NSAIDs, ED₃₀ values were used for comparison. All treatment groups were coded to eliminate observer bias. With this protocol, the ED₃₀ for Indomethacin is 1.0 mg/kg. Representative results are shown in Table IV.

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TABLE III*

	Example	Whole Cells			Microsomes		
		Conc. (nM)	COX-2 % inhib.	COX-1 % inhib.	Conc. (nM)	COX-2 % inhib.	COX-1 % inhib.
5	1	100	96	12	100	53	8
	2	10	69	0	10	49	25
	3	10	42		10	33	19
10	3	100	100		100	76	12
	4				10	47	2
	5	10	0	0	10	43	31
	6	100	78		100	19	16
	7	100	74	0	1000	58	16
15	8	10	41				
	8	100	89				
	9	100	83		100	37	9
	10	100	95		100	71	12
	11	100	39		100	46	7
20	12	100	54				
	13	10	41		10	52	7
	13	100	84		10	58	10
	14	10	73		10	45	29
	14	100	89		100	63	0
25	14	1000	101		1000	69	0

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Example	Conc. (nM)	COX-2 % inhib.	COX-1 % inhib.	Conc. (nM)	COX-2 % inhib.	COX-1 % inhib.
15	20	39				
15	80	76				
15	160	95				
16	20	41				
16	40	50				
16	160	85				
17	40	41				
17	160	77				
18	40	24				
18	160	58				
19	40	21				
19	160	59				
20	10	70				
20	40	91				
21	10	50				
21	40	94				
22	20	39				
22	160	98				
23	20	50				
23	160	88				
24	40	43				
24	160	78				
25	160	40				
26	80	27				
26	160	39				
27	20	38				
27	160	97				

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Example	Conc. (nM)	COX-2 % inhib.	COX-1 % inhib.	Conc. (nM)	COX-2 % inhib.	COX-1 % inhib.
28	20	48				
28	160	69				
29	20	78				
29	160	85				
30	160	30				
31	20	49				
31	160	87				
32	5	43				
32	10	73				
32	40	92				
32	80	99				
33	160	6				
34	10	30				
34	40	80				
34	160	102				
35	20	32				
35	40	57				
35	160	83				
36	10	11				
36	40	50				
36	160	89				
37	10	53				
37	40	82				
37	160	93				
38	10	25				
38	40	63				
38	160	88				
39	10	17				

- 85 -

Example	Conc. (nM)	COX-2 % inhib.	COX-1 % inhib.	Conc. (nM)	COX-2 % inhib.	COX-1 % inhib.
39	160	84				
40	10	43				
40	40	72				
40	160	96				
41						
41						
42	20	10				
42	160	44				
43	10	78				
43	40	101				
44	20	14				
44	40	55				
44	160	106				
45	10	16				
45	40	61				
45	160	101				
46	10	76				
46	40	94				
46	160	97				
47	10	61				
47	40	74				
47	160	101				
48	10	7				
48	160	47				
49	10	53				
49	40	91				
49	80	99				
50	80	42				

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Example	Conc. (nM)	COX-2 % inhib.	COX-1 % inhib.	Conc. (nM)	COX-2 % inhib.	COX-1 % inhib.
51	5	49				
51	20	95				
51	40	102				
52	10	50				
52	40	82				
52	160	102				
53	10	54				
53	40	96				
53	160	102				
54	10	81				
54	80	91				
54	160	99				
55	10	48				
55	80	59				
55	160	65				

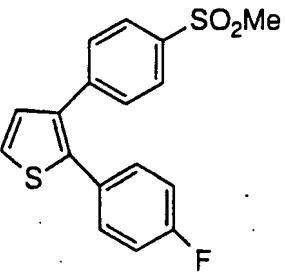
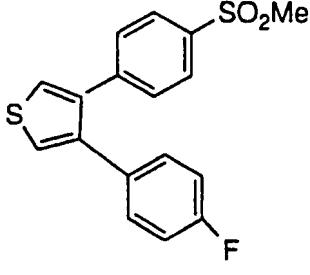
* In the whole cell assay Ibuprofen has an IC50 for COX-1 of 1000 nM, and an IC50 for COX-2 of 3000 nM. Similarly, Indomethacin has an IC50 for COX-1 of 100 nM, and an IC50 for COX-2 of 10 nM.

25

30

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TABLE IV

<u>ED30(mg/kg)</u>	<u>STRUCTURE</u>
5 ~3.00	
10 >10.00	

25

30

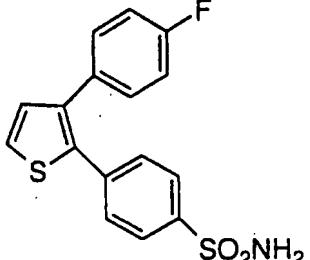
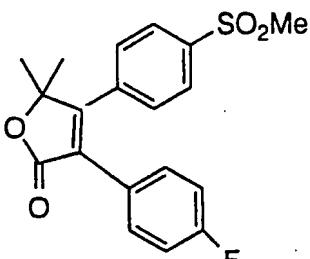
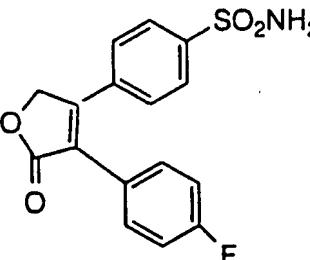
- 88 -

1.40	
5 10 15 20 25	2.80 (in 1% methocel) 0.72
30	0.43

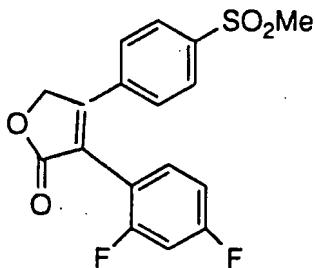
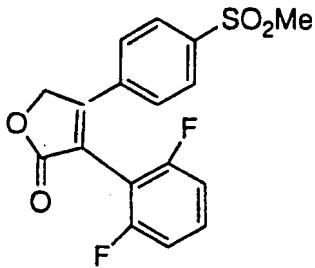
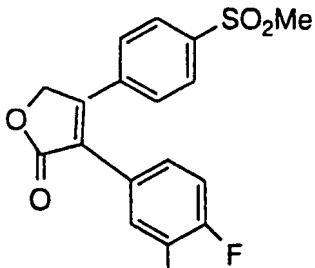
- 89 -

~ 3.00	<chem>CC(C)(C)C1=CSC2=C1C(F)=CC=C2S(=O)(=O)N</chem>
>3.00 3.00	<chem>CC1=CSC2=C1C(F)=CC=C2S(=O)(=O)N</chem>
1.10	<chem>CC1=CSC2=C1C(F)=CC=C2S(=O)(=O)C</chem>

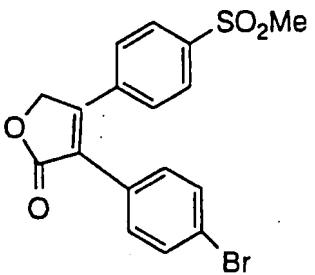
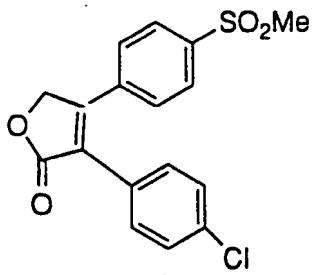
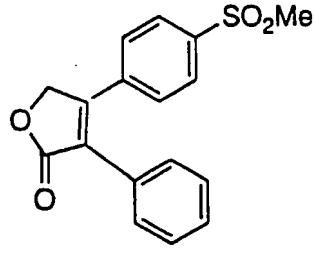
- 90 -

<0.30	 Chemical structure: 2-(4-fluorophenyl)-3-(4-(sulfonyl)phenyl)thiophene. It consists of a thiophene ring fused to a benzene ring, which is substituted at the 4-position with a 4-fluorophenyl group and at the 3-position with a 4-(sulfonyl)phenyl group.
0.42	 Chemical structure: 2-(4-fluorophenyl)-3-(4-(methylsulfonyl)phenyl)thiophene. It consists of a thiophene ring fused to a benzene ring, which is substituted at the 4-position with a 4-(methylsulfonyl)phenyl group and at the 3-position with a 4-fluorophenyl group.
0.034	 Chemical structure: 2-(4-fluorophenyl)-3-(4-(sulfonyl)phenyl)thiophene. It consists of a thiophene ring fused to a benzene ring, which is substituted at the 4-position with a 4-(sulfonyl)phenyl group and at the 3-position with a 4-fluorophenyl group.

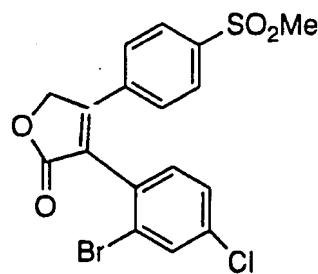
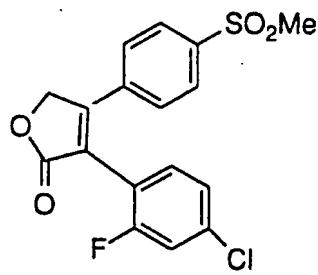
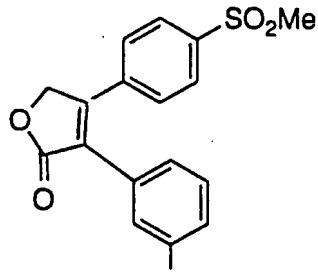
- 91 -

2.03	
1.49	
0.35	

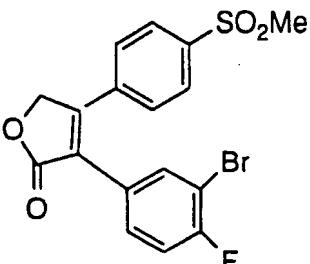
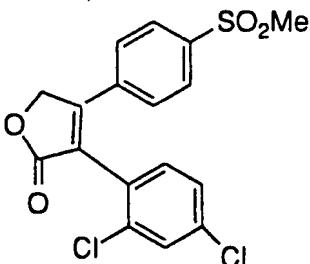
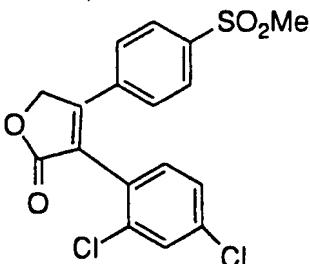
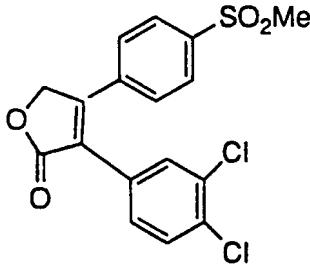
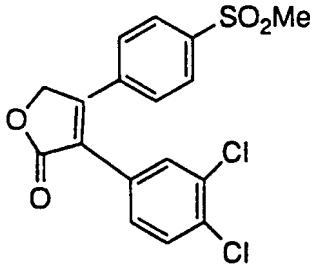
- 92 -

5	0.33	
10	0.90	
15	0.38	

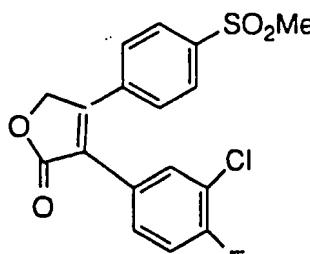
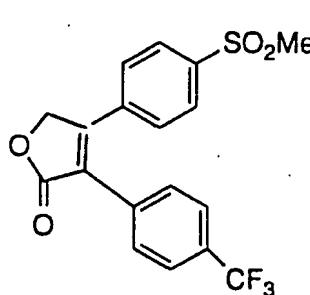
- 93 -

5	0.88	
10	0.47	
15	0.71	

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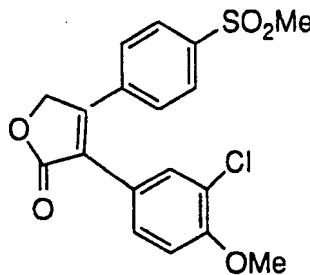
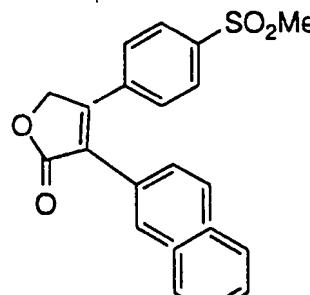
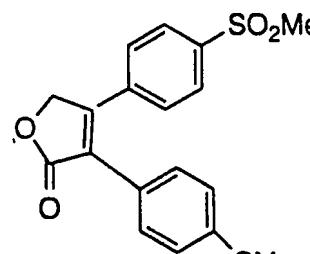
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5	1.85	
10	1.85	
15	0.22 0.23	
20	0.22 0.23	
25		

- 95 -

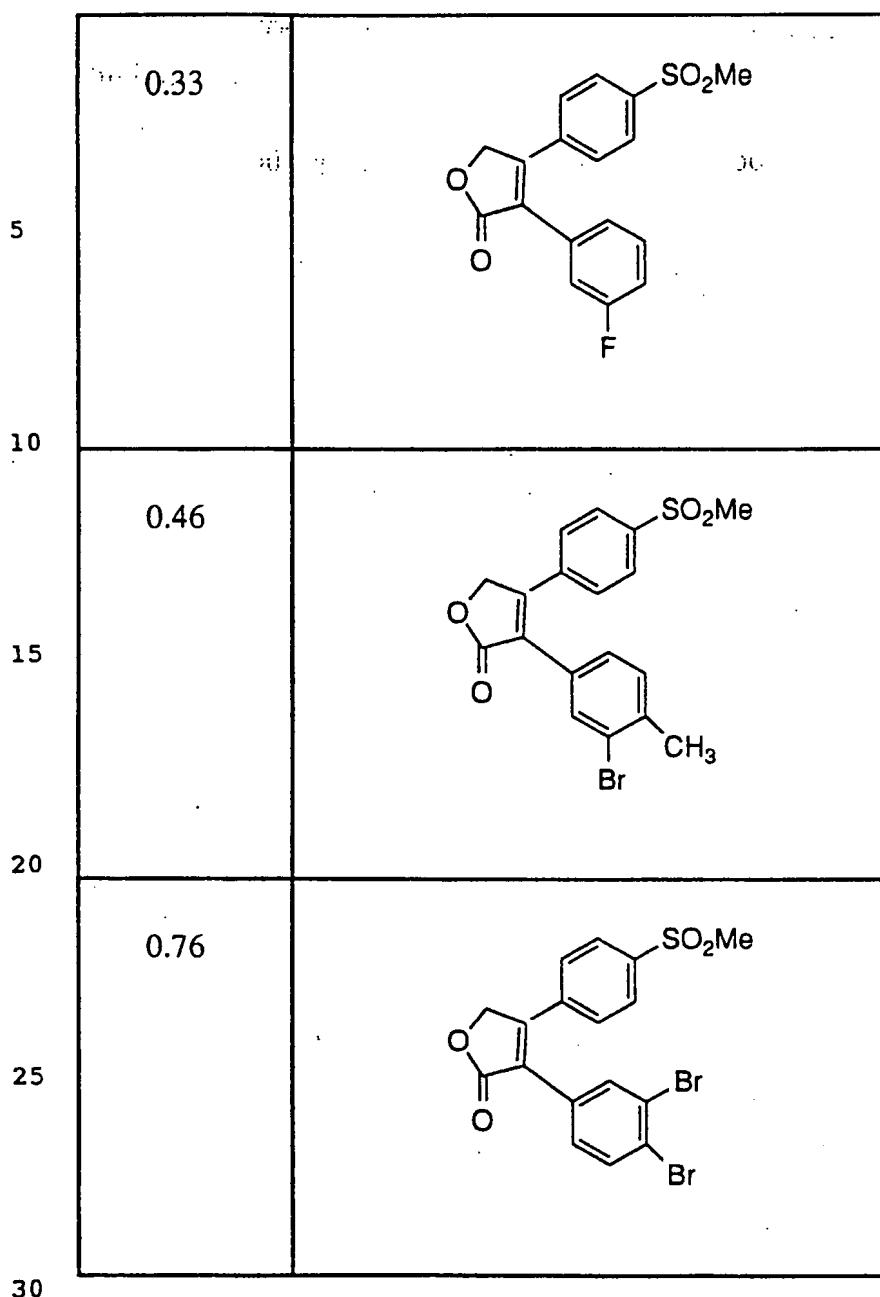
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15	0.81	

30

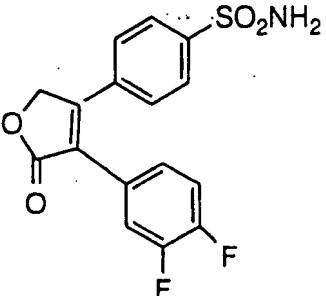
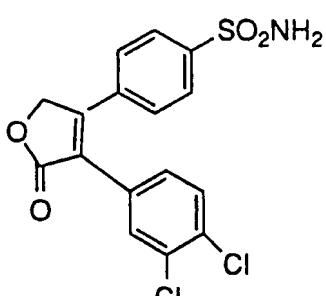
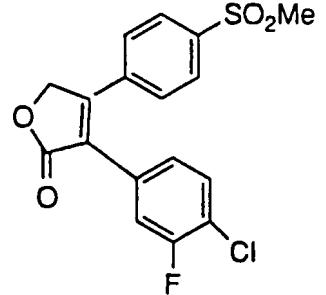
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5	0.68	
10	0.16	
15		
20	~1.00	
25		

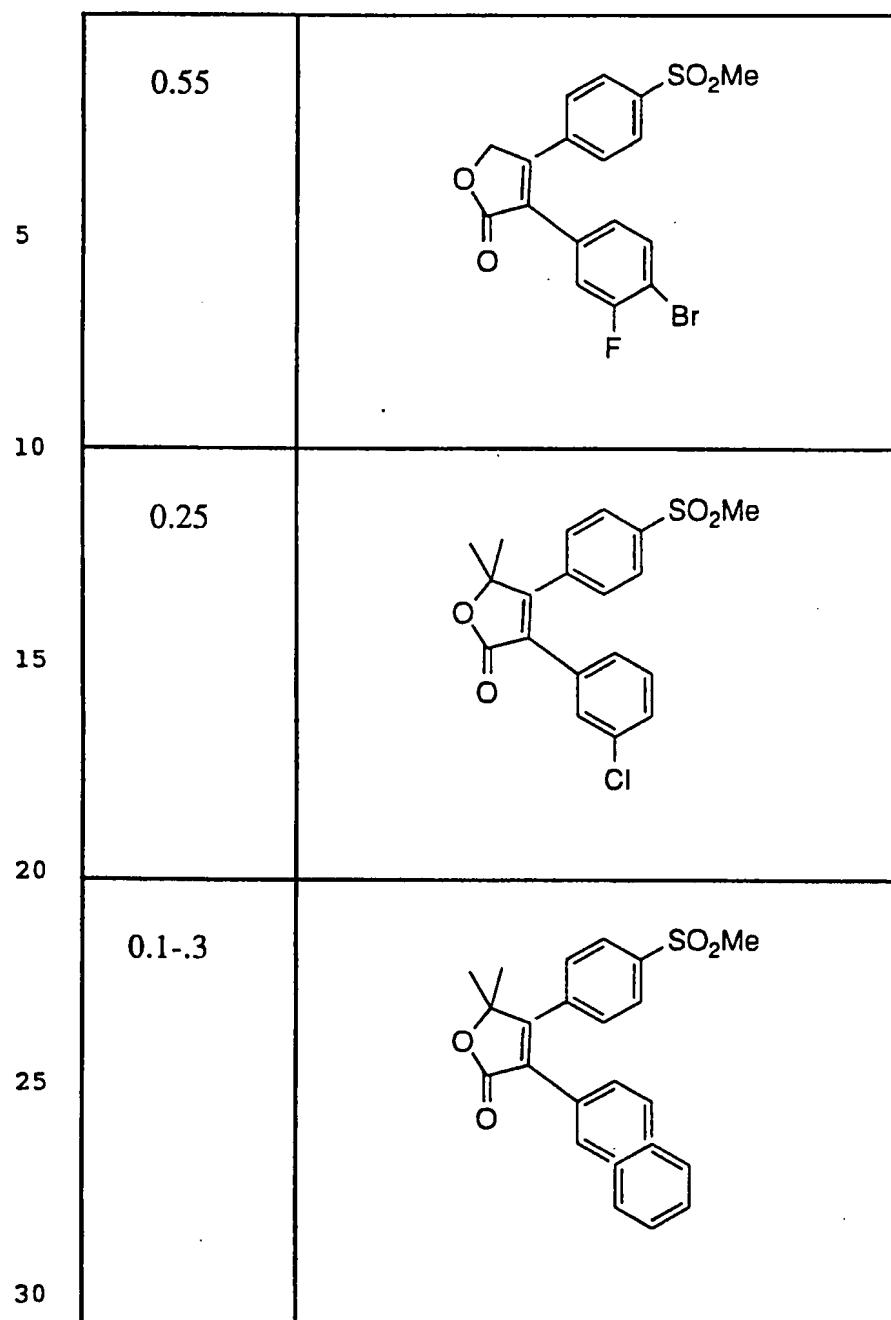
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0.48	
0.46	
0.26	

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~0.10	
0.13	
0.07	

The invention will now be illustrated by the following non-limiting examples in which, unless stated otherwise:

(i) all operations were carried out at room or ambient temperature, that is, at a temperature in the range 18-25°C; evaporation of solvent was carried out using a rotary evaporator under reduced pressure (600-4000 pascals: 4.5-30 mm. Hg) with a bath temperature of up to 60°C; the course of reactions was followed by thin layer chromatography (TLC) and reaction times are given for illustration only; melting points are uncorrected and 'd' indicates decomposition; the melting points given are those obtained for the materials prepared as described; polymorphism may result in isolation of materials with different melting points in some preparations; the structure and purity of all final products were assured by at least one of the following techniques: TLC, mass spectrometry, nuclear magnetic resonance (NMR) spectrometry or microanalytical data; yields are given for illustration only; when given, NMR data is in the form of delta (δ) values for major diagnostic protons, given in parts per million (ppm) relative to tetramethylsilane (TMS) as internal standard, determined at 300 MHz or 400 MHz using the indicated solvent; conventional abbreviations used for signal shape are: s. singlet; d. doublet; t. triplet; m. multiplet; br. broad; etc.: in addition "Ar" signifies an aromatic signal; chemical symbols have their usual meanings; the following abbreviations have also been used v (volume), w (weight), b.p. (boiling point), m.p. (melting point), L (liter(s)), mL (milliliters), g (gram(s)), mg (milligrams(s)), mol (moles), mmol (millimoles), eq (equivalent(s)).

25

The following abbreviations have the indicated meanings:

Ac	=	acetyl
Bn	=	benzyl
DBU	=	1,8-diazabicyclo[5.4.0]undec-7-ene
30		
DIBAL	=	diisobutylaluminum hydride
DMAP	=	4-(dimethylamino)pyridine

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	DMF	=	N,N-dimethylformamide
	Et ₃ N	=	triethylamine
	LDA	=	lithium diisopropylamide
	m-CPBA	=	metachloroperbenzoic acid
5	MMPP	=	monoperoxyphthalic acid
	MPPM	=	monoperoxyphthalic acid, magnesium salt 6H ₂ O
	Ms	=	methanesulfonyl = mesyl = SO ₂ Me
	MsO	=	methanesulfonate = mesylate
10	NSAID	=	non-steroidal anti-inflammatory drug
	OXONE®	=	2KHSO ₅ •KHSO ₄ •K ₂ SO ₄
	PCC	=	pyridinium chlorochromate
	PDC	=	pyridinium dichromate
	Ph	=	phenyl
15	Phe	=	benzenediyl
	Pye	=	pyridinediyl
	r.t.	=	room temperature
	rac.	=	racemic
	SAM	=	aminosulfonyl or sulfonamide or SO ₂ NH ₂
20	TBAF	=	tetra-n-butylammonium fluoride
	Th	=	2- or 3-thienyl
	TFAA	=	trifluoroacetic acid anhydride
	THF	=	tetrahydrofuran
	Thi	=	thiophenediyl
25	TLC	=	thin layer chromatography
	TMS-CN	=	trimethylsilyl cyanide
	Tz	=	1H (or 2H)-tetrazol-5-yl
	C ₃ H ₅	=	allyl
30	<u>Alkyl Group Abbreviations</u>		
	Me	=	methyl

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	Et	=	ethyl
	n-Pr	=	normal propyl
	i-Pr	=	isopropyl
	n-Bu	=	normal butyl
	i-Bu	=	isobutyl
5	s-Bu	=	secondary butyl
	t-Bu	=	tertiary butyl
	c-Pr	=	cyclopropyl
	c-Bu	=	cyclobutyl
	c-Pen	=	cyclopentyl
10	c-Hex	=	cyclohexyl

15

20

25

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EXAMPLE 1

3-(4-Aminosulfonyl)phenyl)-2-(4-fluorophenyl)-5-(2-hydroxy-2-propyl)thiophene

5 Step 1: 1-(4-Fluorophenyl)-2-(4-(methylthio)phenyl)ethanone
To 4-fluorobenzaldehyde (5.40 g) in 1,2-dichloroethane (43.50 mL) were added TMS-CN (4.32 g) and ZnI₂ (44 mg). After 0.5 h at r.t., the solvent was removed *in vacuo*. To the resulting TMS cyanohydrin (9.20 g) in THF (42.0 mL) at -78°C was added dropwise a
10 solution of LDA 0.51M in THF (88.9 mL). After a period of 0.5 h, a THF solution (30.0 mL) of 4-(chloromethyl)thioanisole (9.93 g) was added dropwise over 0.5 h. After 18 h at +5°C, the resulting mixture was treated with TBAF (57.5 mL) followed by a 25% aqueous solution of NH₄OAc (100 mL) and extracted with EtOAc (2 x 150 mL). After
15 evaporation, a 10:1 mixture of Et₂O and hexane (200 mL) was added to the crude ketone. After stirring for 10 h and filtration, the title product was obtained as a solid by filtration (2.40 g).
1H NMR (CD₃COCD₃): δ 2.45 (3H, s), 4.34 (2H, s), 7.19-7.29 (6H, m),
8.14 (2H, q).

20 Step 2: Cis,trans-3-chloro-3-(4-fluorophenyl)-2-(4-(methylthio)phenyl)propenal
To a solution of 1-(4-fluorophenyl)-2-(4-(methylthio)phenyl)ethanone (2.50 g) in 1,2-dichloroethane (27.0 mL) were introduced the
25 Vilsmeier reagent (Aldrich catalog, 1992-1993) 3.3M (11.6 mL) and DMAP (1.17 g). After a period of 4 h at 80°C, the reaction mixture was extracted with EtOAc and 25% aqueous solution of NH₄OAc. After evaporation *in vacuo* and drying for a few hours, the title product was used as such for the next step.
30 1H NMR (CD₃COCD₃): δ 2.40 and 2.48 (3H, 2s), 6.90-7.80 (8H, m), 9.55 (1H, s).

Step 3: 5-(4-Fluorophenyl)-4-(4-(methylthio)phenyl)thiophene-2-carboxylic acid methyl ester

To a solution of *cis,trans* 3-chloro-3-(4-fluorophenyl)-2-(4-(methylthio)phenyl)propenal (3.00 g) in pyridine (12.0 mL) were added 5 methyl thioglycolate (1.16 mL) and Et₃N (4.09 mL). The resulting mixture was then heated at 80°C for 2 h. After extraction with EtOAc and washing with 3N HCl, the title product was purified by flash chromatography (30% EtOAc in hexane) (2.00 g).

10 ¹H NMR (CD₃COCD₃): δ 2.48 (3H, s), 3.88 (3H, s), 7.11 (2H, t), 7.21 (4H, s), 7.37 (2H, q), 7.80 (1H, s).

Step 4: 5-(4-Fluorophenyl)-4-(4-(methylsulfinyl)phenyl)thiophene-2-carboxylic acid methyl ester

To a solution of 5-(4-fluorophenyl)-4-(4-(methylthio)phenyl)-15 thiophene-2-carboxylic acid methyl ester (5.60 g) in CH₂Cl₂ (84.0 mL) at 0°C was added portionwise m-CPBA 50 to 60% (5.39 g). After TLC showed completion (50% EtOAc in hexane), the reaction mixture was extracted with saturated NaHCO₃, dried over Na₂SO₄, filtered and evaporated to dryness to provide the title compound as a white foam (5.00 20 g).

10 ¹H NMR (CD₃COCD₃): δ 2.75 (3H, s), 3.92 (3H, s), 7.15 (2H, t), 7.40 (2H, q), 7.52 (2H, d), 7.66 (2H, d), 7.90 (1H, s).

Step 5: 4-(4-(Aminosulfonyl)phenyl)-5-(4-fluorophenyl)thiophene-2-carboxylic acid methyl ester

5-(4-Fluorophenyl)-4-(4-(methylsulfinyl)phenyl)thiophene-2-carboxylic acid methyl ester (0.500 g) was dissolved in TFAA (10.0 mL) and refluxed for 0.5 h. The solvent was then removed *in vacuo* and the resulting residue was co-evaporated 10 times with a Et₃N-MeOH solution 30 (1:1) (100.0 mL) to provide a viscous oil after pumping for a few hours. The oil was dissolved in HOAc (10.0 mL) and treated at +10°C with Cl₂ in

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HOAc (1.9M) (3.5 mL). After 20 min., the solvent was removed under reduced pressure and after pumping, THF (20.0 mL) was added to the resulting mass of product. After bubbling NH₃ through for a few minutes at 0°C, the reaction mixture was stirred for 0.5 h at r.t. After extraction with EtOAc - 25% NH₄OAc solution and flash chromatography (30 to 5 40% EtOAc in hexane), the title product was obtained as a white solid (0.210 g).

¹H NMR (CD₃COCD₃): δ 3.90 (3H, s), 6.55 (2H, bs), 7.13 (2H, t), 7.40 (2H, q), 7.46 (2H, d), 7.83 (2H, d), 7.90 (1H, s).

10 Step 6: 3-(4-Aminosulfonyl)phenyl)-2-(4-fluorophenyl)-5-(2-hydroxy-2-propyl)thiophene

To 4-(4-aminosulfonyl)phenyl)-5-(4-fluorophenyl)thiophene-2-carboxylic acid methyl ester (0.460 g) in THF (5.70 mL) at 0°C was added MeMgBr (1.4M) in toluene-THF solution (5.00 mL). The mixture 15 was then stirred at r.t. for a few hours. The reaction was quenched by the addition of 25% NH₄OAc solution, extracted with EtOAc and dried over Na₂SO₄. The title compound was purified by flash chromatography (40 to 50% EtOAc in hexane) (0.300 g).

1H NMR (CD₃COCD₃): δ 1.65 (6H, s), 4.52 (1H, s), 6.55 (2H, bs), 7.09 (20 3H, m), 7.34 (2H, dd), 7.30 (2H, m), 7.43 (2H, d), 7.82 (2H, d). Anal. calcd. for C₁₉H₁₈FNO₃S₂; C, 58.31; H, 4.60; N, 3.58. Found: C, 57.94; H, 4.66; N, 3.44

EXAMPLE 2

25

3-(4-(Aminosulfonyl)phenyl)-2-(4-fluorophenyl)thiophene

Step 1: 4-(4-(Aminosulfonyl)phenyl)-5-(4-fluorophenyl)thiophene-2-carboxylic acid

30

To a solution of 4-(4-(aminosulfonyl)phenyl)-5-(4-fluorophenyl)thiophene-2-carboxylic acid methyl ester (Example 1, Step 5)

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(0.210 g) in THF (2.0 mL) were added MeOH (1.0 mL), NaOH 1N (1.0 mL) and a few drops of NaOH 10N. The resulting mixture was heated at 45°C for 2 h and the reaction was then partitioned between EtOAc and HCl (3N) to provide the title product as a white solid (0.200 g).

1H NMR (CD₃COCD₃) δ 6.60 (2H, s), 7.15 (2H, t), 7.35 (2H, q), 7.45 (2H, d), 7.82 (2H, d), 7.87 (1H, s).

Step 2: 3-(4-(Aminosulfonyl)phenyl)-2-(4-fluorophenyl)thiophene

To a solution of 3-(4-(aminosulfonyl)phenyl)-2-(4-fluorophenyl)thiophene-2-carboxylic acid (0.280 g) in quinoline (4.0 mL) was added Cu bronze (0.300 g). After 0.5 h at 180°C under nitrogen, the reaction mixture was extracted with EtOAc and HCl 3N, dried over Na₂SO₄ and purified by flash chromatography (30% EtOAc in hexane) to give the title compound as a white solid (0.180 g).

1H NMR (CD₃COCD₃): δ 6.60 (2H, bs), 7.15 (2H, t), 7.29 (1H, d), 7.35 (2H, q), 7.45 (2H, d), 7.60 (1H, d), 7.83 (2H, d).

Anal. calcd for C₁₆H₁₂FNO₂S₂:

C, 57.65; H, 3.60; N, 4.20.

Found: C, 57.62; H, 3.59; N, 4.15.

20

EXAMPLE 3

3-(4-(Aminosulfonyl)phenyl)-2-(4-fluorophenyl)-5-(2-propyl)thiophene

25 1H NMR (CD₃COCD₃) δ 1.40 (6H, d), 3.25 (1H, septuplet), 6.58 (2H, bs), 7.05 (1H, s), 7.15 (2H, t), 7.32 (2H, dd), 7.46 (2H, d), 7.80 (2H, d).

Anal. calcd. for C₁₉H₁₈FNO₂S₂.

C, 60.80; H, 4.80; N, 3.73.

30 Found: C, 60.59; H, 4.45; N, 3.60.

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EXAMPLE 4

3-(4-(Aminosulfonyl)phenyl)-2-cyclohexylthiophene

1H NMR (CD₃)₂CO) δ 1.24-1.40 (3H, m), 1.40-1.56 (2H, m), 1.65-1.85
5 (3H, m), 1.90-2.0 (2H, m), 3.18 (1H, m), 6.58 (2H, bs), 7.05 (1H, d), 7.37
(1H, d), 7.58 (2H, d), 7.97 (2H, d).

EXAMPLE 5

10 5-(4-Carboxyphenyl)-4-(4-(methylsulfonyl)phenyl)thiophene-2-carboxylic
acid

Step 1: 4-(2-(4-Methylthiophenyl)-1-oxo-ethyl)benzoic acid methyl
ester

15 To methyl 4-formylbenzoate (10.30 g) in 1,2-dichloroethane
at r.t. were added TMS-CN (6.58 mL) and ZnI₂ (2.00 g), after 0.5 h at
r.t., the solvent was removed *in vacuo*. To the resulting TMS cyanohyrin
(5.00 g) in THF (22.0 mL) at -78°C was added dropwise a solution of LDA
0.87 M in THF (26.2 mL). After a period of 0.5 h, a THF solution (10.0
20 mL) of 4-(chloromethyl)thioanisole was added dropwise over 0.5 h. The
temperature was then brought slowly to -20°C then to 5°C for 2 h and
TBAF 1M in THF (50.0 mL) was added. After the addition of 25%
aqueous solution of NH₄OAc, the reaction mixture was extracted with
EtOAc, dried over NaSO₄, evaporated *in vacuo* and purified by flash
25 chromatography (20 to 30% EtOAc in hexane) to afford the title
compound as a white solid (7.00 g).

Step 2: 4-(1-Oxo-2-(4-(methylsulfonyl)phenyl)ethyl) benzoic acid
methyl ester

30 To 7.10 g of 4-(2-(4-methylthiophenyl)-1-oxo-ethyl)benzoic
acid methyl ester in MeOH (100 mL) was added oxone (21.0 g) in H₂O

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(20.0 mL) at 0°C. After a few hours at r.t., the reaction mixture was extracted with EtOAc and H₂O to afford after flash chromatography (50 to 100% EtOAc in hexane), the title product as a white solid (3.20 g).
1H NMR (CD₃COCD₃) δ 3.10 (3H, s), 3.95 (3H, s), 4.65 (2H, s), 7.60 (2H, d), 7.96 (2H, d), 8.20 (4H, q).

5

Step 3: Cis,trans 4-(1-Chloro-3-oxo-2-(4-(methylsulfonyl)phenyl)-1-propenyl)benzoic acid methyl ester

To a solution of 4-(1-oxo-2-((4-methylsulfonyl)phenyl)ethyl) benzoic acid (1.70 g) in 1,2-dichloroethane (15.0 mL) were added the 10 Vilsmeier reagent 3.3 M (6.2 mL) and DMAP (0.624 g). The resulting mixture was heated at 80°C for 4 h. The reaction mixture was then extracted with 25% aqueous solution of NH₄OAc and EtOAc. After drying over Na₂SO₄ and evaporation the title compound was obtained as an oil and used as such for the next step.

15

Step 4: 5-(4-(Methoxycarbonyl)phenyl)-4-(4-(methylsulfonyl)-phenyl)thiophene-2-carboxylic acid methyl ester

Prepared from 4-(1-chloro-3-oxo-2-(4-methylsulfonyl)-phenyl)-1-propenyl)benzoic acid methyl ester as for Example 1, Step 3.
20 1H NMR (CD₃COCD₃) δ 3.13 (3H, s), 3.85 and 3.92 (6H, 2s), 7.50 (2H, d), 7.55 (2H, d), 7.90 (2H, d), 7.92 (1H, s), 7.92 (2H, d).

25

Step 5: 5-(4-(Carboxyphenyl)-4-(4-(methylsulfonyl)phenyl)-thiophene-2-carboxylic acid

Prepared from 5-(4-(methoxycarbonyl)phenyl)-4-(4-(methylsulfonyl)phenyl) thiophene-2-carboxylic acid methyl ester as for Example 2, Step 1.
1H NMR (CD₃COCD₃) δ 3.15 (3H, s), 7.50 (2H, d), 7.62 (2H, d), 7.95 (2H, d), 7.98 (1H, s), 8.05 (2H, d).

30

Anal calcd. for C₁₉H₁₄O₆S₂•0.1 H₂O:

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 C, 56.46; H, 3.51.
 Found: C, 56.18; H, 3.51.

EXAMPLE 6

5 4-(4-Fluorophenyl)-2-methyl-5-(4-(methylsulfonyl)phenyl)thiazole

Step 1: 1-(4-Fluorophenyl)-2-(4-(methylsulfonyl)phenyl)ethanone

To 1-(4-Fluorophenyl)-2-(4-(methylthio)phenyl)ethanone of Example 1, Step 1 (17.9 g) in a solution of CH₂Cl₂-MeOH (272.0 mL/27.0 mL) at 0°C was added MPPM (28.0 g). The cooling bath was then removed and the reaction mixture stirred at r.t. for 1 h. At 0°C, additional MPPM (28.0 g) was added and the reaction mixture kept for 1.5 h at r.t. The insoluble material was filtered followed by evaporation of the solvents, the residue was then extracted with CH₂Cl₂-NaHCO₃. After evaporation *in vacuo*, the resulting solid was washed with ether-hexane (1:1) and filtered to provide the title compound 16.8 g.
1H NMR (CD₃COCD₃) δ 3.13 (3H, s), 3.58 (2H, s), 7.29 (2H, t), 7.55 (2H, d), 7.88 (2H, d), 8.20 (2H, dd).

20 Step 2: 2-Bromo-1-(4-fluorophenyl)-2-(4-(methylsulfonyl)phenyl)-ethanone

To 1-(4-Fluorophenyl)-2-(4-(methylsulfonyl)phenyl)ethanone (1.00 g) in CH₂Cl₂ containing CHCl₃ (1.0 mL) and CCl₄ (1.0 mL) was added bromine (0.614 g). After shining light for 1 h, the reaction was quenched with Na₂S₂O₄, extracted with CH₂Cl₂, dried over Na₂SO₄ and evaporated to yield the title compound which was used as such for the next step (1.10 g).
1H NMR (CD₃COCD₃) δ 3.10 (3H, s), 7.05 (1H, s), 7.30 (2H, t), 7.87 (2H, d), 7.95 (2H, d), 8.25 (2H, dd).

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Step 3: 4-(4-Fluorophenyl)-2-methyl-5-(4-(methylsulfonyl)phenyl)-thiazole

To 2-bromo-1-(4-fluorophenyl)-2-(4-(methylsulfonyl)phenyl)-ethanone (1.10 g) in ethanol (15.0 mL) were added thioacetamide (0.266 g) and pyridine (0.300 mL). After refluxing for 2 h, the reaction mixture
5 was extracted with EtOAc, 25% NH₄OAc and purified by flash chromatography (50% EtOAc in hexane then 90% Et₂O in hexane) to yield the title compound (0.320 g).

10 ¹H NMR (CD₃COCD₃) δ 2.72 (3H, s), 3.15 (3H, s), 7.09 (2H, t), 7.52 (2H, dd), 7.60 (2H, d), 7.92 (2H, d).

15 Anal. calcd. for C₁₇H₁₄FNO₂S₂:

C, 58.78; H, 4.03; N, 4.03.

Found: C, 58.71, H, 4.17; N, 3.85.

EXAMPLE 7

15

2-(4-Fluorophenyl)-3-(4-(methylsulfonyl)phenyl)-2-cyclopenten-1-one

Step 1: 1-(4-Fluorophenyl)-5-hexen-2-one

To a suspension of 14.6 g (80 mmol) of CdCl₂ in 200 mL of ether cooled to 0°C was added 115 mL of 1.3 M solution of 3-butene-1-magnesium bromide dropwise. The mixture was refluxed for 1 h and ether was then removed by distillation. Benzene (500 mL) was introduced, followed by a solution of 17.5 g (100 mmol) 4-fluorophenylacetyl chloride. After refluxing for 1 h, the reaction mixture was quenched with 200 mL of saturated aqueous NH₄Cl, 50 mL of 1 N HCl, and extracted with 200 mL of 1:1 hexane/EtOAc. The organic phase was dried over MgSO₄ and concentrated. The residue was purified by flash chromatography eluted with 4:1 hexane/EtOAc to give 15 g of the title product.

30 ¹H NMR (CDCl₃) δ 2.40 (2H, t), 2.53 (2H, t), 3.63 (2H, s), 4.90-4.98 (2H, m), 5.67-5.78 (1H, m), 6.98 (2H, t), 7.13 (2H, m).

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Step 2: 1-(4-Fluorophenyl)-5-oxo-2-pentanone

A solution of 14 g of 1-(4-fluorophenyl)-5-hexen-2-one in 200 mL of 3:1 CH₂Cl₂/MeOH was cooled to -78°C and treated with excess ozone. The resulting mixture was treated with 15 g of triphenylphosphine and stirred at room temperature for 1 h. The reaction mixture was concentrated and flash chromatographed with 3:1 hexane/EtOAc to give 8 g of the title ketoaldehyde.

⁵ ¹⁰ ^{1H} NMR (CDCl₃) δ 2.72 (4H, s), 3.71 (2H, s), 6.99 (2H, t), 7.14 (2H, m), 9.73 (1H, s).

Step 3: 2-(4-Fluorophenyl)-2-cyclopenten-1-one

A solution of 8 g of 1-(4-fluorophenyl)-5-oxo-2-pentanone in 300 mL of MeOH was treated with 2 g of NaOMe. The mixture was stirred for 2 h and then quenched with 5 mL of HOAc. The solvent was ¹⁵ evaporated and the residue purified by flash chromatography, eluting with 3:1 hexane/EtOAc to give 7 g of the title product.

¹⁰ ¹⁵ ^{1H} NMR (CDCl₃) δ 2.57 (2H, m), 2.68 (2H, m), 7.04 (2H, J=8.8 Hz, t), 7.67 (2H, J=8.8, 5.5 Hz, dd), 7.77 (1H, m).

²⁰ **Step 4: 1-(4-(Methylthio)phenyl)-2-(4-fluorophenyl)-2-cyclopenten-1-ol**

To a solution of 3.86 g (19 mmol) of 4-bromothioanisole in 90 mL of Et₂O cooled at -78°C, was added 22 mL of 1.7 M solution of t-BuLi in pentane (38 mmol) dropwise. The reaction mixture was stirred for 15 ²⁵ min at -78°C and a solution of 2.23 g of 2-(4-Fluorophenyl)-2-cyclopenten-1-one in 10 mL of Et₂O was added. After stirring for 15 min at -78°C, the reaction mixture was warmed to 0°C, and quenched with 50 mL of sat. NH₄Cl. The product was extracted with 100 mL EtOAc, dried over Na₂SO₄, and purified by flash chromatography, eluted with 4:1 ³⁰ hexane/EtOAc to give 3.4 g of the desired product.

¹H NMR (CDCl₃) δ 2.12 (1H, s), 2.34 (2H, m), 2.44 (3H, s), 2.45-2.52 (1H, m), 2.56-2.65 (1H, m), 6.37 (1H, m), 6.84 (2H, J=8.7 Hz, t), 7.17 (2H, J=8.3 Hz, d), 7.24-7.33 (4H, m).

⁵ Step 5: 2-(4-Fluorophenyl)-3-(4-(methylthio)phenyl)-2-cyclo-
penten-1-one

To a suspension of PCC (4.5 g, 20.9 mmol) and 10 g of anhydrous 4Å molecular sieves in 150 mL of CH₂Cl₂ was added a solution of 2.2 g (7.3 mmol) of 1-(4-(methylthio)phenyl)-2-(4-fluorophenyl)-2-cyclopenten-1-ol in 20 mL CH₂Cl₂. The mixture was stirred for 1 h at r.t. and then diluted with 300 mL of Et₂O. After filtration and concentration, the residue was flash chromatographed with 2:1 hexane/EtOAc to give 1.5 g of the title product.

¹⁰ ¹⁵ ¹H NMR (CDCl₃) δ 2.45 (3H, s), 2.68 (2H, m), 3.00 (2H, m), 7.02 (2H, J=8.6 Hz, t), 7.11 (2H, J=8.6 Hz, d), 7.15-7.23 (4H, m).

¹⁵ Step 6: 2-(4-Fluorophenyl)-3-(4-(methylsulfonyl)phenyl)-2-
cyclopenten-1-one

To a solution of 50 mg (0.17 mmol) of 2-(4-Fluorophenyl)-3-(4-methylthio)phenyl)-2-cyclopenten-1-one in 8 mL of 10:1 CH₂Cl₂/MeOH was added 124 mg (0.2 mmol) of MPPM. The reaction mixture was stirred at room temperature for 2 h and then diluted with 10 mL of 1:1 hexane/EtOAc. After filtration and concentration, the residue was purified by flash chromatography eluted with 2:1 EtOAc/hexane to give 45 mg of the title product.

²⁰ ²⁵ ¹H NMR (acetone-d₆) δ 2.67 (2H, m), 3.14 (3H, s), 3.16 (2H, m), 7.05-7.10 (2H, m), 7.20-7.25 (2H, m), 7.63 (2H, d), 7.93 (2H, d).

EXAMPLE 8

³⁰ 4-(4-(Methylsulfonyl)phenyl)-5-(4-fluorophenyl)-isothiazole

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To a solution of 338 mg (1 mmol) of cis,trans 3-chloro-3-(4-fluorophenyl)-2-(4-(methylsulfonyl)phenyl)propenal in 5 mL of acetone was added 230 mg (3 mmol) of NH₄SCN. The reaction mixture was refluxed for 3 h, and then quenched with 20 mL of saturated NaHCO₃. The product was extracted with 100 mL of EtOAc, dried over Na₂SO₄, 5 concentrated and purified by flash chromatography eluted with 3:2 hexane/EtOAc to give 250 mg of the title product.
1H NMR (CDCl₃) δ 8.57 (1H, s), 7.93 (3H, d), 7.50 (2H, d), 7.30 (2H, t), 7.08 (2H, t).

10

EXAMPLE 9

3-(4-Fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Step 1: 2-Bromo-1-(4-(methylsulfonyl)phenyl)ethanone

15 A solution of 197 g of 4-(Methylthio)acetophenone (ref: JACS, 1952, 74, p. 5475) in 700 mL of MeOH and 3500mL of CH₂Cl₂ was added 881 g of MMPP over a period of 30 min. After 3 h at room temperature the reaction mixture was filtered and the filtrate was washed with 2 L of saturated aqueous solution of NaHCO₃ and 1 L of brine. The 20 aqueous phase was further extracted with 2 L of CH₂Cl₂. The combined extracts was dried over Na₂SO₄ concentrated to give 240 g of 4-(methylsulfonyl)acetophenone as a white solid.

To a cooled (-5 °C) solution of 174 g of 4-(methylsulfonyl)acetophenone in 2.5 L of CHCl₃ was added 20 mg of 25 AlCl₃, followed by a solution of 40 mL of Br₂ in 300 mL CHCl₃. The reaction mixture was then treated with 1.5 L of water and the CHCl₃ was separated. The aqueous layer was extracted with 1 L of EtOAc. The combined extracts was dried over Na₂SO₄ and concentrated. The crude product was recrystallized from 50/50 EtOAc/hexane to give 210 g of 30 2-bromo-1-(4-(methylsulfonyl)phenyl)ethanone as a white solid.

Step 2:

To the product of Step 1 (216 mg) dissolved in acetonitrile (4 mL) was added Et₃N (0.26 mL), followed by 4-fluorophenylacetic acid (102 mg). After 1.5 h at room temperature 0.23 mL of DBU was added. The reaction mixture was stirred for another 45 min and then treated with 5 mL of 1N HCl. The product was extracted with EtOAc, dried over Na₂SO₄ and concentrated. The residue was purified by flash chromatography (40% EtOAc in hexane) to yield 150 mg of the title compound as a solid.

10 ¹H NMR (CD₃COCD₃) δ 3.15 (3H, s), 5.36 (3H, s), 7.18 (2H, J=8.9 Hz, t), 7.46 (2H, m), 7.7 (2H, J=8.65 Hz, d), 7.97 (2H, J=8.68, d).

EXAMPLE 103-(4-Fluorophenyl)-4-(4-(aminosulfonyl)phenyl)-2-(2H)-furanone

15 ¹H NMR (CD₃COCD₃) δ 5.34 (2H, s), 6.67 (2H, bd), 7.18 (2H, m), 7.46 (2H, m), 7.61 (2H, m), 7.90 (2H, m).
M.P. 187-188 °C (d).

20

EXAMPLE 113-(4-Fluorophenyl)-4-(4-(methylsulfonyl)phenyl)furanStep 1:

25 Using the product of Example 10, (0.2 g) in THF (5 mL) and toluene (3 mL) was added slowly at -78°C a solution of DIBAL (0.72 mL, 1M in toluene). After 15 min, the solution was warmed up to 0°C for another 15 min. This mixture was then poured into a chilled aqueous solution of sodium potassium tartrate and EtOAc. The organic layer was 30 stirred for 0.5 h with a few crystals of camphor sulfonic acid. This

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solution was then concentrated and purified by flash chromatography to yield the title compound.

^1H NMR (CDCl₃) δ 3.1 (3H, s), 7.02 (2H, J=8.9, t), 7.18 (2H, m), 7.4 (2H, J=8.8 Hz, d), 7.58 (1H, s), 7.68 (1H, s), 7.85 (2H, J=8.8 Hz, d)

5

EXAMPLE 12

5,5-Dimethyl-3-(4-fluorophenyl)-4-(4-methylsulfonylphenyl)-2-(5H)-furanone

10 **Step 1:** Methyl 2-trimethylsilyloxyisobutyrate

To a solution of 1.2 mL (10.4 mmol) of methyl 2-hydroxyisobutyrate in 50 mL of CH₂Cl₂ were added 1.2 g (17.6 mmol) of imidazole and 2.1 mL (16.6 mmol) of TMSCl. The mixture was stirred at r.t. for 1.5 h and quenched with 20 mL of H₂O. The organic layer was dried over MgSO₄, concentrated and passed through a short plug of silica gel eluted with 9:1 hexane/EtOAc. Evaporation of solvent afforded 1.27 g of the title compound as a colorless oil.

^1H NMR (CD₃COCD₃) δ 0.08 (9H, s), 1.38 (6H, s), 3.67 (3H, s).

20 **Step 2:** 2-Trimethylsilyloxy-4'-(methylthio)isobutyrophenone

A solution of 204 mg (1.0 mmol) of 4-bromothioanisole in 2.5 mL of THF was cooled to -78°C and treated with 0.42 mL of 2.5 M n-BuLi solution in hexane. After stirring at -78°C for 1 h, a solution of 380 mg (2.0 mmol) of methyl 2-trimethylsilyloxyisobutyrate in 2 mL of THF was added. The mixture was stirred at -78°C for 2 h and then quenched with NH₄OAc buffer. The product was extracted with EtOAc, dried over MgSO₄ and concentrated. The residue was purified by flash chromatography, eluting with 19:1 hexane/EtOAc to give 95 mg of the title product.

30 ^1H NMR (CD₃COCD₃) δ 0.05 (9H, s), 1.52 (6H, s), 2.53 (3H, s), 7.33 (2H, d), 8.12 (2H, d).

Step 3: 2-Hydroxy-4'-(methylthio)isobutyrophenone

To a solution of 40 mg (0.14 mmol) of 2-trimethylsilyloxy-4'-(methylthio)isobutyrophenone in 2 mL THF was added 0.2 mL of 1 M n-Bu4NF in THF. The resulting mixture was stirred for 30 min and then 5 quenched with 10 mL of NH4OAc buffer. The product was extracted with EtOAc, dried over MgSO4 and concentrated. The residue was purified by flash chromatography, eluting with 4:1 hexane/EtOAc to give 25 mg of the title product.

10 ^1H NMR (CD3COCD3) δ 1.50 (6H, s), 2.54 (3H, s), 4.68 (1H, s), 7.30 (2H, d), 8.15 (2H, d).

Step 4: 2-(4-Fluorophenylacetoxy)-4'-(methylthio)isobutyrophenone

To a solution of 72 mg (0.34 mmol) 2-hydroxy-4'-(methylthio)isobutyrophenone in 1.7 mL of CH2Cl2 were added 0.2 mL of 15 pyridine and 140 mg (0.81 mmol) of 4-fluorophenylacetyl chloride. The mixture was stirred at room temperature overnight and then quenched with NH4OAc buffer. The product was extracted with EtOAc, dried over MgSO4 and concentrated. The crude product was purified by flash chromatography eluting with 8:1 hexane/EtOAc to give 95 mg of the title 20 product.

10 ^1H NMR (CD3COCD3) δ 1.62 (3H, s), 1.67 (3H, s), 2.48 (3H, s), 3.79 (2H, s), 7.0-7.3 (6H, m), 7.78 (2H, d).

Step 5: 5,5-Dimethyl-3-(4-fluorophenyl-4-(4-methylthiophenyl)-2-(5H)-furanone

To a solution of 95 mg of 2-(4-fluorophenylacetoxy)-4'-(methylthio)-isobutyrophenone in 4 mL of CH2Cl2 was added 0.2 mL of 1,8-diazabicyclo(5.4.0)undec-7-ene. The mixture was stirred for 4 h and 30 diluted with NH4OAc buffer. The product was extracted with EtOAc, dried over MgSO4 and concentrated. The residue was purified by flash

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chromatography, eluting with 20:1 toluene/EtOAc to give 75 mg of the title product.

1H NMR (CD₃COCD₃) δ 1.58 (6H, s), 2.50 (3H, s), 7.03 (2H, dd), 7.25-7.35 (4H, m), 7.41 (2H, dd).

5 Step 6: 5,5-Dimethyl-3-(4-fluorophenyl)-4-(4-methylsulfonylphenyl)-2-(5H)-furanone

To a solution of 81 mg of 5,5-dimethyl-3-(4-fluorophenyl)-4-(4-methyl-thiophenyl)-2-oxo-2H-dihydrofuran in 1.8 mL of CH₂Cl₂ and 0.2 mL of MeOH was added 250 mg of MPPM. The reaction mixture was 10 stirred at room temperature for 1 h and then quenched with aqueous NaHCO₃. The product was extracted with EtOAc, dried over MgSO₄ and concentrated. The crude product was purified by flash chromatography eluting with 1:1 hexane/EtOAc to give 73 mg of the title product.
1H NMR (CD₃COCD₃) δ 1.62 (6H, s), 3.15 (3H, s), 7.02 (2H, dd), 7.40 (2H, dd), 7.65 (2H, d), 8.03 (2H, d).

EXAMPLE 13

2-((4-aminosulfonyl)phenyl)-3-(4-fluorophenyl)thiophene

20 1H NMR (CD₃COCD₃) δ 6.60 (2H, bs), 7.12 (2H, t), 7.25 (1H, d), 7.35 (2H, m), 7.45 (2H, d), 7.65 (1H, d), 7.85 (2H, d).

25 Analysis calculated for C₁₆H₁₂FNS₂O₂
C, 57.65; H, 3.60; N, 4.20
Found: C, 57.55; H, 3.79; N, 4.03

EXAMPLE 14

3-(4-(Trifluoroacetylaminosulfonyl)phenyl)-2-(4-fluorophenyl)thiophene

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¹H NMR (300 MHz, CD₃COCD₃) δ 7.15 (2H, t), 7.30 (3H, m), 7.45 (2H, d), 7.65 (1H, d), 7.95 (2H, d).

EXAMPLE 15

5

3-(2,4-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₇H₁₂F₂O₄S
C, 58.28; H, 3.45; S, 9.15
10 Found: C, 58.27; H, 3.50; S, 9.27

EXAMPLE 16

15 3-(3,4-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone
To a solution of 3,4-difluorophenylacetic acid (ALDRICH
CHIMICAL) (10 g) and 2-bromo-1-(4-(methylsulfonyl)phenyl)ethanone (Example 9, Step 1) (17.3 g) in acetonitrile (200 mL) at room temperature was added slowly triethylamine (20.2 mL). After 1 h at room
temperature, the mixture was cooled in an ice bath and treated with 17.4
20 mL of DBU. After 2 h at 0°C, the mixture was treated with 200 mL of 1N
HCl and the product was extracted with EtOAc, dried over Na₂SO₄ and
concentrated. The residue was applied on top of a silica gel plug (sintered
glass funnel) eluted with 75% EtOAc/hexane, giving after evaporation of
the solvent and swish in ethyl acetate, 10 g of the title compound.

25

Analysis calculated for C₁₇H₁₂F₂O₄S
C, 58.28; H, 3.45; S, 9.15
Found: C, 58.02; H, 3.51; S, 9.35

30

EXAMPLE 17

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3-(2,6-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₇H₁₂F₂O₄S
 C, 58.28; H, 3.45; S, 9.15
Found: C, 58.18; H, 3.50; S, 9.44

5

EXAMPLE 18

3-(2,5-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

10 Analysis calculated for C₁₇H₁₂F₂O₄S
 C, 58.28; H, 3.45; S, 9.15
Found: C, 58.89; H, 3.51; S, 9.11

15

EXAMPLE 19

3-(3,5-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₇H₁₂F₂O₄S
 C, 58.28; H, 3.45; S, 9.15
20 Found: C, 58.27; H, 3.62; S, 9.32

EXAMPLE 20

3-(4-Bromophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

25 Analysis calculated for C₁₇H₁₃BrO₄S
 C, 51.94; H, 3.33; S, 8.16
Found: C, 51.76; H, 3.42; S, 8.21

30

EXAMPLE 21

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3-(4-Chlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

¹H NMR (300 MHz, CDCl₃) δ 7.93 (2H, d), 7.49 (2H, d), 7.35 (4H, m), 5.16 (2H, s), 3.06 (3H, s)

5

EXAMPLE 22

3-(4-Methoxyphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₈H₁₆O₅S
10 C, 62.78 H, 4.68; S, 9.31
Found: C, 62.75; H, 4.72; S, 9.39

EXAMPLE 23

15 3-(Phenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

To a solution of phenylacetic acid (27.4 g, 201 mmol) and 2-bromo-1-(4-(methylsulfonyl)phenyl)ethanone (Example 9, Step 1) (60 g, 216 mmol, 1.075 eq.) in acetonitrile (630 mL) at 25°C was added slowly triethylamine (30.8 mL, 1.1 eq.). The mixture was stirred for 20 min. at 20 room temperature and then cooled in an ice bath. DBU (60.1 mL, 3 eq.) was slowly added. After stirring for 20 min. in the ice bath, the reaction was complete and the mixture was acidified with 1N HCl (color changes from dark brown to yellow). Then 2.4 L of ice and water were added, stirred for a few minutes, then the precipitate was filtered and rinsed with 25 water (giving 64 g of crude wet product). The solid was dissolved in 750 mL of dichloromethane (dried over MgSO₄, filtered) and 300 g of silica gel was added. The solvent was evaporated to near dryness (silica gel a bit sticky) and the residue was applied on top of a silica gel plug (sintered glass funnel) eluted with 10% EtOAc/CH₂Cl₂, giving after evaporation of the 30 solvent and swish in ethyl acetate, 36.6 g (58%) of the title compound.

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Analysis calculated for C₁₇H₁₄O₄S
C, 64.95; H, 4.49; S, 10.20
Found: C, 64.63; H, 4.65; S, 10.44

EXAMPLE 24

5

3-(2-Chlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₇H₁₃ClO₄S
C, 58.54; H, 3.76; S, 9.19
10 Found: C, 58.59; H, 3.80; S, 9.37

EXAMPLE 25

15 3-(2-Bromo-4-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₇H₁₂BrFO₄S
C, 49.75; H, 2.93
20 Found: C, 49.75; H, 3.01

EXAMPLE 26

25 3-(2-Bromo-4-Chlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

1H NMR (300 MHz, acetone-d₆) δ 7.95 (2H, d), 7.85 (1H, d), 7.63 (2H, dd), 7.55 (1H, dd), 7.45 (1H, d), 5.50 (2H, s), 3.15 (3H, s)

EXAMPLE 27

30

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3-(4-Chloro-2-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

¹H NMR (300 MHz, acetone-d₆) δ 8.0 (2H, d), 7.70 (2H, d), 7.50-7.30 (3H, m), 5.35 (2H, s), 3.15 (3H, s)

5

EXAMPLE 28

3-(3-Bromo-4-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

10

Analysis calculated for C₁₇H₁₂BrFO₄S

C, 49.75; H, 2.93

Found: C, 49.44; H, 2.98

15

EXAMPLE 29

3-(3-Chlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

20

Analysis calculated for C₁₇H₁₃ClO₄S

C, 58.54; H, 3.76

Found: C, 58.29; H, 3.76

EXAMPLE 30

25

3-(2-Chloro-4-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

30

Analysis calculated for C₁₇H₁₂ClFO₄S

C, 55.67; H, 3.30

Found: C, 55.67; H, 3.26

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EXAMPLE 31

3-(2,4-Dichlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₇H₁₂Cl₂O₄S
5 C, 53.28; H, 3.16; S, 8.37
 Found: C, 52.89; H, 3.23; S, 8.58

EXAMPLE 32

10 3-(3,4-Dichlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₇H₁₂Cl₂O₄S
 C, 53.28; H, 3.16; S, 8.37
15 Found: C, 53.07; H, 3.32; S, 8.51

EXAMPLE 33

3-(2,6-Dichlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

20 Analysis calculated for C₁₇H₁₂Cl₂O₄S
 C, 53.28; H, 3.16; S, 8.37
 Found: C, 52.99; H, 3.22; S, 8.54

EXAMPLE 34

25 3-(3-Chloro-4-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

30 ¹H NMR (300 MHz, acetone-d₆) d 8.0 (2H, d), 7.70 (2H, d), 7.60 (1H, d),
 7.25-7.40 (2H, m), 5.35 (2H, s), 3.15 (3H, s)

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EXAMPLE 35

3-(4-Trifluoromethylphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

5 ^1H NMR (CD₃COCD₃) δ 8.10 (2H, d), 7.82-7.93 (4H, m), 7.75 (2H, d),
5.55 (2H, s), 3.30 (3H, s)

EXAMPLE 36

10 3-(3-Fluoro-4-methoxyphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₈H₁₅FO₅S
C, 59.66; H, 4.17

15 Found: C, 59.92; H, 4.37

EXAMPLE 37

20 3-(3-Chloro-4-methoxyphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₈H₁₅ClO₅S
C, 57.07; H, 3.99

25 Found: C, 57.29; H, 4.15

EXAMPLE 38

30 3-(3-Bromo-4-methoxyphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₈H₁₅BrO₅S

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 C, 51.08; H, 3.57
Found: C, 51.38; H, 3.62

EXAMPLE 39

5 3-(2-Fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₇H₁₃FO₄S
 C, 61.44; H, 3.94
Found: C, 61.13; H, 3.85

10

EXAMPLE 40

3-(4-Methylthiophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

15 ¹H NMR (300 MHz, acetone-d₆) d 8.0 (2H, d), 7.70 (2H, d), 7.35 (2H, d),
7.25 (2H, d), 5.35 (2H, s), 3.15 (3H, s), 2.55 (3H, s)

EXAMPLE 41

20 3-(3-Fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

¹H NMR (300 MHz, CDCl₃) d 7.93 (2H, d), 7.49 (2H, d), 7.35 (1H, m),
7.12 (3H, m), 5.18 (2H, s), 3.06 (3H, s)

25

EXAMPLE 42

3-(2-Chloro-6-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

30 ¹H NMR (300 MHz, acetone-d₆), d 8.0 (2H, d), 7.70 (2H, d), 7.55-7.65
(1H, m), 7.40 (1H, d), 7.30 (1H, m), 5.60 (2H, s), 3.15 (3H, s)

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EXAMPLE 43

3-(3-Bromo-4-methylphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

5

Analysis calculated for C₁₈H₁₅BrO₄S
 C, 53.08; H, 3.71
Found: C, 53.06; H, 3.83

10

EXAMPLE 44

3-(4-Bromo-2-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

15

Analysis calculated for C₁₇H₁₂BrFO₄S
 C, 49.65; H, 2.94
Found: C, 49.76; H, 3.00

20

EXAMPLE 45

3-(3,4-Dibromophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

¹H NMR (300 MHz, acetone-d₆) δ 8.0 (2H, d), 7.80 (1H, d), 7.75 (3H, m),
7.25 (1H, d), 5.35 (2H, s), 3.15 (sH, s)

25

EXAMPLE 46

3-(4-Chloro-3-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

30

Analysis calculated for C₁₇H₁₂ClFO₄S

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 C, 55.67; H, 3.30
Found: C, 55.45; H, 3.30

EXAMPLE 47

5 3-(4-Bromo-3-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₇H₁₂BrFO₄S
 C, 49.66; H, 2.94; S, 7.80
10 Found: C, 49.79; H, 3.01; S, 7.51

EXAMPLE 48

15 3-(4-Bromo-2-chlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Analysis calculated for C₁₇H₁₂BrClO₄S
 C, 47.74; H, 2.83; S, 7.50
20 Found: C, 47.92; H, 2.84; S, 7.42

EXAMPLE 49

3-(2-Naphthyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

25 Analysis calculated for C₂₁H₁₆O₄S
 C, 69.22; H, 4.43
Found: C, 69.22; H, 4.46

EXAMPLE 50

30 3-(7-Quinoliny)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

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Analysis calculated for C₂₀H₁₅NO₄S
 C, 65.74; H, 4.14; N, 3.83
Found: C, 65.34; H, 4.40; N, 3.80
M.S. (DCI, CH₄) calculated for M⁺, 365
5 Found for M⁺⁺¹, 366

EXAMPLE 51

10 3-(3,4-Dichlorophenyl)-4-(4-(aminosulfonyl)phenyl)-2-(2H)-furanone
1H NMR (400 MHz, CD₃COCD₃) δ 7.92 (2H, dd), 7.64 (3H, dm), 7.60
(1H, dd), 7.32 (1H, dd), 6.70 (1H, bs), 5.38 (2H, s)

EXAMPLE 52

15 3-(3,4-Difluorophenyl)-4-(4-(aminosulfonyl)phenyl)-2-(2H)-furanone
1H NMR (400 MHz, CD₃COCD₃) δ 7.92 (2H, dd), 7.64 (2H, dd), 7.30-
7.45 (2H, m), 7.22 (1H, m), 6.68 (2H, bs), 5.37 (2H, s)
20

EXAMPLE 53

25 3-(3-Chloro-4-methoxyphenyl)-4-(4-(aminosulfonyl)phenyl)-2-(2H)-furanone
Analysis calculated for C₁₇H₁₄ClNO₅S
 C, 53.76; H, 3.72, N, 3.69
Found: C, 53.32; H, 3.84, N, 3.59
M.S. (DCI, CH₄) calculated for M⁺, 379
30 Found for M⁺⁺¹, 380

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EXAMPLE 54

5 3-(3-Bromo-4-methoxyphenyl)-4-(4-(aminosulfonyl)phenyl)-2-(2H)-furanone

Analysis calculated for C₁₇H₁₄BrNO₅S
 C, 48.13; H, 3.33, N, 3.30
Found: C, 48.26; H, 3.40, N, 3.28
10 M.S. (DCI, CH₄) calculated for M⁺, 423
 Found for M⁺⁺¹, 424

EXAMPLE 55

15 3-(Phenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

20 Into a 20 ml glass ampule are added 1 g of 2-(4-(methylsulfonyl)phenyl)phenylacetylene, 20 mg of Rh₄(CO)₁₂, 1.5 g of Et₃N, 10 ml of THF, 1 ml of water under nitrogen atmosphere, and the ampule is placed in a 100-ml stainless steel autoclave. The reaction system is flushed three times with CO then charged at room temperature to a initial CO pressure of 100 atm. The reaction is carried at 100 °C for 5 h. The solution is then diluted with 50 ml of benzene and washed with brine, 1N HCl. The benzene solution is dried over Na₂SO₄, and concentrated. The crude products are separated by column chromatography on silica gel 25 eluted with 2:1 EtOAc/hexane to give the title compound and its regioisomer.

EXAMPLE 56

30 3-(Phenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

Step 1: 2-trimethylsilyloxy-4-(4-(methylthio)phenyl)-3,4-dihydrofuran

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To a solution of 3.86 g (19 mmol) of 4-bromothioanisole in 90 mL of Et₂O cooled at -78°C, is added 22 mL of 1.7 M solution of t-BuLi in pentane (38 mmol) dropwise. The reaction mixture is stirred for 15 min at -78°C and 3.8 g of CuI is added and the reaction mixture is allowed to warm to -40 °C over a period of 30 min. A solution of 1.7 g of 2(5H)-

5 furanone in 10 ml of THF is added. After stirring for 1 h, 2 ml of freshly distilled TMSCl is added dropwise. The reaction mixture is then treated with 2 ml of Et₃N and 50 ml of sat. NaHCO₃, and extracted with 100 ml of ether. The ether layer is dried over Na₂SO₄ and concentrated to the crude title compound which is used for the next step without further purification.

10

Step 2: 4-(4-(methylthio)phenyl)-2-(5H)-furanone

To a solution of 4 g of Pd(OAc)₂ in 100 ml of acetonitrile is added dropwise the crude product from Step 1(5 g) under nitrogen at room

15 temperature. After 10 h at room temperature, the mixture is condensed under reduced pressure and the residue is purified by flash chromatography on silica gel eluted with 2:1 hexane/EtOAc to give the title compound.

20

Step 3: 3-iodo-4-(4-(methylthio)phenyl)-2-(5H)-furanone

To a solution of 3 g of the product of Step 2 in 30 ml of pyridine is added 8.7 g of I₂. The mixture is stirred for 24 h and then diluted with 200 ml of ether, washed with 100 ml of 5N HCl and 50 ml of

25 5N Na₂S₂O₃. The ether layer is dried over Na₂SO₄ and concentrated to give the title compound.

Step 4: 3-(Phenyl)-4-(4-(methylthio)phenyl)-2-(5H)-furanone

30

A mixture of 4 g of the product of Step 3, 3.7 g of PhB(OH)₂, 0.4 g of Ph₃As, 0.4 g of PdCl₂(PhCN)₂ in 100 ml of benzene and 15 ml of

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2N NaOH is refluxed for 6 h. Ether(200 ml) is then added and the mixture is washed with 100 ml of saturated NaHCO₃. The organic layer is dried over MgSO₄ and concentrated. The residue is purified by flash chromatography on silica gel eluted with 4:1 hexane/EtOAc to give the title compound.

5

Step 5: 3-(Phenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone

To a solution of 3 g of the product of Step 4 in 80 mL of 10:1 CH₂Cl₂/MeOH is added 5.5 g of MPPM. The reaction mixture is stirred at 10 room temperature for 2 h and then diluted with 100 mL of 1:1 hexane/EtOAc. After filtration and concentration, the residue is purified by flash chromatography eluted with 2:1 EtOAc/hexane to give the title product.

15

20

25

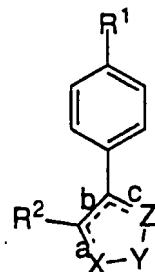
30

WHAT IS CLAIMED IS:

1. A compound of formula I

5

10



I

or a pharmaceutically acceptable salt thereof wherein:

X-Y-Z-is selected from the group consisting of:

- 15 (a) -CH₂CH₂CH₂-,
- (b) -C(O)CH₂CH₂-,
- (c) -CH₂CH₂C(O)-,
- (d) -CR⁵(R^{5'})-O-C(O)-,
- (e) -C(O)-O-CR⁵(R^{5'})-,
- 20 (f) -CH₂-NR³-CH₂-,
- (g) -CR⁵(R^{5'})-NR³-C(O)-,
- (h) -CR⁴=CR^{4'}-S-,
- (i) -S-CR⁴=CR^{4'}-,
- (j) -S-N=CH-,
- 25 (k) -CH=N-S-,
- (l) -N=CR⁴-O-,
- (m) -O-CR⁴=N-
- (n) -N=CR⁴-NH-,
- (o) -N=CR⁴-S-, and
- 30 (p) -S-CR⁴=N-,
- (q) -C(O)-NR³-CR⁵(R^{5'})-,

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(r) $-\text{NR}^3-\text{CH}=\text{CH}-$ provided R^1 is other than $-\text{S}(\text{O})_2\text{Me}$,
(s) $-\text{CH}=\text{CH}-\text{NR}^3-$ provided R^1 is other than $-\text{S}(\text{O})_2\text{Me}$,

when side b is a double bond, and sides a and c are single bonds; and

5 X-Y-Z-is selected from the group consisting of:

(a) $=\text{CH}-\text{O}-\text{CH}=$, and
(b) $=\text{CH}-\text{NR}^3-\text{CH}=$,
(c) $=\text{N}-\text{S}-\text{CH}=$,
10 (d) $=\text{CH}-\text{S}-\text{N}=$,
(e) $=\text{N}-\text{O}-\text{CH}=$,
(f) $=\text{CH}-\text{O}-\text{N}=$,
(g) $=\text{N}-\text{S}-\text{N}=$,
15 (h) $=\text{N}-\text{O}-\text{N}=$,

when sides a and c are double bonds and side b is a single bond;

15 R^1 is selected from the group consisting of

(a) $\text{S}(\text{O})_2\text{CH}_3$,
(b) $\text{S}(\text{O})_2\text{NH}_2$,
(c) $\text{S}(\text{O})_2\text{NHC}(\text{O})\text{CF}_3$,
20 (d) $\text{S}(\text{O})(\text{NH})\text{CH}_3$,
(e) $\text{S}(\text{O})(\text{NH})\text{NH}_2$,
(f) $\text{S}(\text{O})(\text{NH})\text{NHC}(\text{O})\text{CF}_3$,
(g) $\text{P}(\text{O})(\text{CH}_3)\text{OH}$, and
25 (h) $\text{P}(\text{O})(\text{CH}_3)\text{NH}_2$,

R^2 is selected from the group consisting of

25 (a) C₁₋₆alkyl,
(b) C₃, C₄, C₅, C₆, and C₇, cycloalkyl,
(c) mono-, di- or tri-substituted phenyl wherein the substituent is
30 selected from the group consisting of
(1) hydrogen,
(2) halo,
(3) C₁₋₆alkoxy,

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(4) C₁₋₆alkylthio,
(5) CN,
(6) CF₃,
(7) C₁₋₆alkyl,
(8) N₃,
5 (9) -CO₂H,
(10) -CO₂-C₁₋₄alkyl,
(11) -C(R⁵)(R⁶)-OH,
(12) -C(R⁵)(R⁶)-O-C₁₋₄alkyl, and
(13) -C₁₋₆alkyl-CO₂-R⁵;

10 (d) mono-, di- or tri-substituted heteroaryl wherein the heteroaryl is a monocyclic aromatic ring of 5 atoms, said ring having one hetero atom which is S, O, or N, and optionally 1, 2, or 3 additionally N atoms; or
15 the heteroaryl is a monocyclic ring of 6 atoms, said ring having one hetero atom which is N, and optionally 1, 2, 3, or 4 additional N atoms; said substituents are selected from the group consisting of
20 (1) hydrogen,
(2) halo, including fluoro, chloro, bromo and iodo,
(3) C₁₋₆alkyl,
(4) C₁₋₆alkoxy,
(5) C₁₋₆alkylthio,
(6) CN,
25 (7) CF₃,
(8) N₃,
(9) -C(R⁵)(R⁶)-OH,
(10) -C(R⁵)(R⁶)-O-C₁₋₄alkyl;

R³ is selected from the group consisting of
30 (a) hydrogen,
(b) CF₃,
(c) CN,

(d) C₁-6alkyl,
(e) hydroxyC₁-6alkyl, and
(f) -C(O)-C₁-6alkyl,
(g) optionally substituted
(1) -C₁-5 alkyl-Q,
(2) -C₁-3alkyl-O-C₁-3alkyl-Q,
(3) -C₁-3alkyl-S-C₁-3alkyl-Q,
(4) -C₁-5 alkyl-O-Q, or
(5) -C₁-5 alkyl-S-Q,
wherein the substituent resides on the alkyl and the substituent
is C₁-3alkyl,
(h) -Q,
R⁴ and R^{4'} are each independently selected from the group consisting of
(a) hydrogen,
(b) CF₃,
(c) CN,
(d) C₁-6alkyl,
(e) -Q,
(f) -O-Q;
(g) -S-Q, and
(h) optionally substituted
(1) -C₁-5 alkyl-Q,
(2) -O-C₁-5 alkyl-Q,
(3) -S-C₁-5 alkyl-Q,
(4) -C₁-3alkyl-O-C₁-3alkyl-Q,
(5) -C₁-3alkyl-S-C₁-3alkyl-Q,
(6) -C₁-5 alkyl-O-Q,
(7) -C₁-5 alkyl-S-Q,
wherein the substituent resides on the alkyl and the substituent
is C₁-3alkyl, and
R⁵, R^{5'} and R⁶, R⁷ and R⁸ are each independently selected from the
group consisting of

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- (a) hydrogen,
- (b) C₁-6alkyl,

or R⁵ and R⁶ or R⁷ and R⁸ together with the carbon to which they are attached form a monocyclic saturated carbon ring of 3, 4, 5, 6 or 7 atoms;

5

Q is CO₂H, CO₂-C₁-4alkyl, tetrazolyl-5-yl, C(R⁷)(R⁸)(OH), or C(R⁷)(R⁸)(O-C₁-4alkyl),

10 provided that when X-Y-Z is -S-CR⁴=CR^{4'}, then R⁴ and R^{4'} are other than CF₃.

2. A compound according to Claim 1 wherein X-Y-Z-is selected from the group consisting of:

- (a) -CH₂CH₂CH₂-,
- (b) -C(O)CH₂CH₂-,
- (c) -CH₂CH₂C(O)-,
- (d) -CR⁵(R^{5'})-O-C(O)-,
- (e) -C(O)-O-CR⁵(R^{5'})-,
- (f) -CH₂-NR³-CH₂-,
- (g) -CR⁵(R^{5'})-NR³-C(O)-,
- (h) -CR⁴=CR^{4'}-S-,
- (i) -S-CR⁴=CR^{4'}-,
- (j) -S-N=CH-,
- (k) -CH=N-S-,
- (l) -N=CR⁴-O-,
- (m) -O-CR⁴=N-,
- (n) -N-CR⁴-NH-,
- (o) -N=CR⁴-S-, and
- (p) -S-CR⁴=N-,
- (q) -C(O)-NR³-CR⁵(R^{5'})-,
- (r) -NR³-CH=CH- provided R¹ is other than -S(O)₂Me,

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(s) -CH=CH-NR³. provided R¹ is other than -S(O)2Me,

R¹ is selected from the group consisting of

- (a) S(O)2CH₃,
- (b) S(O)2NH₂,
- (c) S(O)2NHC(O)CF₃,
- 5 (d) S(O)NHCH₃,
- (e) S(O)NHNH₂, and
- (f) S(O)NHNHC(O)CF₃;

R² is selected from the group consisting of

- (a) C₁₋₄alkyl,
- 10 (b) C₃, C₄, C₅, C₆, and C₇, cycloalkyl,
- (c) mono- or di-substituted phenyl wherein the substituent is selected from the group consisting of
 - (1) hydrogen,
 - (2) fluoro, chloro, and bromo,
 - 15 (3) C₁₋₄alkoxy,
 - (4) C₁₋₄alkylthio,
 - (5) CN,
 - (6) CF₃,
 - (7) C₁₋₄alkyl,
 - 20 (8) N₃,
 - (9) -CO₂H,
 - (10) -CO₂-C₁₋₃alkyl,
 - (11) -C(R⁵)(R⁶)-OH, and
 - (12) -C(R⁵)(R⁶)-O-C₁₋₃alkyl,
- 25 (d) mono- or di-substituted heteroaryl selected from the group consisting of
 - (1) furanyl,
 - (2) diazinyl, triazinyl and tetrazinyl,
 - (3) imidazolyl,
 - 30 (4) isooxazolyl,
 - (5) isothiazolyl,

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- (6) oxadiazolyl,
- (7) oxazolyl,
- (8) pyrazolyl,
- (9) pyrrolyl,
- (10) thiadiazolyl,
- 5 (11) thiazolyl,
- (12) thienyl,
- (13) triazolyl, and
- (14) tetrazolyl,

wherein said substituents are selected from the group consisting of

- 10 (a) hydrogen,
- (b) fluoro, chloro, bromo,
- (c) C₁₋₄alkoxy,
- (d) C₁₋₄alkylthio,
- (e) CN,
- 15 (5) CF₃,
- (6) C₁₋₄alkyl,
- (7) N₃,
- (8) -C(R⁵)(R⁶)-OH,
- (9) -C(R⁵)(R⁶)-O-C₁₋₄alkyl.

20

3. A compound according to Claim 2 wherein R² is selected from the group consisting of

- (a) cyclohexyl, and
- (b) mono- or di-substituted phenyl, and

25

wherein the substituents are selected from the group

- consisting of
- (1) hydrogen,
- (2) halo,
- (3) C₁₋₄alkoxy,
- (4) C₁₋₄alkylthio,
- 30 (5) CN,

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- (6) CF₃,
- (7) C₁₋₄alkyl,
- (8) N₃, and
- (9) -C(R⁵)(R⁶)-OH;

R³ is selected from the group consisting of

- 5 (a) hydrogen,
- (b) CF₃,
- (c) C₁₋₃alkyl and hydroxyC₁₋₃alkyl,
- (d) CN,

R⁴ and R^{4'} are each independently selected from the group consisting of

- 10 (a) hydrogen,
- (b) CF₃,
- (c) C₁₋₃alkyl,
- (d) CN,
- (e) chloro and fluoro; and

15 R⁵, R^{5'}, R⁶, are each independently selected from the group consisting of

- (a) hydrogen,
- (b) methyl or ethyl,

or R⁵ and R⁶ together with the carbon to which they are attached
form a saturated carbon ring of 4, 5 or 6 atoms.

20

4. A compound according to claim 3 wherein

X-Y-Z-is selected from the group consisting of:

- (a) -CH₂-O-C(O)-, and
- (b) -C(O)-O-CH₂-, and

25 R¹ is selected from the group consisting of

- (a) S(O)₂CH₃,
- (b) S(O)₂NH₂,
- (c) S(O)NHCH₃, and
- (d) S(O)NHNH₂;

30 R² is

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mono or di-substituted phenyl wherein the substituents are selected from the group consisting of

- (1) hydrogen,
- (2) halo, selected from the group consisting of fluoro, chloro and bromo,
- 5 (3) methoxy, and
- (4) methyl.

5. A compound according to claim 4 wherein X-Y-Z-is selected from the group consisting of:

- 10 (a) -CH₂-O-C(O)-, and
- (b) -C(O)-O-CH₂-, and

R¹ is selected from the group consisting of

- (a) S(O)₂CH₃, and
- (b) S(O)₂NH₂,

15 R² is

mono or di-substituted phenyl wherein the substituents are selected from the group consisting of

- (1) hydrogen,
- (2) halo, selected from the group consisting of fluoro, chloro and bromo.

6. A compound according to Claim 2 wherein R² is a mono- or di-substituted heteroaryl wherein heteroaryl is selected from the group consisting of

- 25 (1) furanyl,
- (2) diazinyl, triazinyl, tetrazinyl,
- (3) imidazolyl,
- (4) isooxazolyl,
- (5) isothiazolyl,
- 30 (6) oxadiazolyl,
- (7) oxazolyl,

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- (8) pyrazolyl,
- (9) pyrrolyl,
- (10) thiadiazolyl,
- (11) thiazolyl,
- (12) thienyl,
- 5 (13) triazolyl, and
- (14) tetrazolyl,

wherein the substituents are selected from the group consisting of

- (a) hydrogen,
- 10 (b) fluoro or chloro,
- (c) C₁-3alkoxy,
- (d) C₁-6alkylthio,
- (e) CN,
- (f) CF₃,
- 15 (g) C₁-3alkyl,
- (h) -C(R⁵)(R⁶)-OH;
- (i) -C(R⁵)(R⁶)-O-C₁-4alkyl.

7. A compound according to Claim 6 wherein
20 the heteroaryl is selected from the group consisting of

- (1) 3-isothiazolyl,
- (2) 4-isothiazolyl,
- (3) 5-isothiazolyl,
- (4) 2-oxazolyl,
- 25 (5) 4-oxazolyl,
- (6) 5-oxazolyl,
- (7) 2-thiazolyl,
- (8) 4-thiazolyl,
- (9) 5-thiazolyl,
- 30 (10) 1,2-diazinyl,
- (11) 1,3-diazinyl, and

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(12) 1,4-diazinyl, and

wherein the substituents are selected from the group consisting of

- (1) hydrogen,
- (2) fluoro or chloro,
- (3) C₁₋₃alkoxy,
- (4) C₁₋₃alkylthio,
- (5) CN,
- (6) C₁₋₃alkyl, and
- (7) -C(R⁵)(R⁶)-OH,

wherein R⁵ and R⁶ are each independently hydrogen, methyl
or ethyl.

10

8. A compound according to claim 7 wherein
X-Y-Z-is selected from the group consisting of:

- (a) -CH₂-O-C(O)-,
- (b) -C(O)-O-CH₂-, and
- (c) -CH₂-NR³-C(O)-;

15

R¹ is selected from the group consisting of

- (a) S(O)₂CH₃,
- (b) S(O)₂NH₂,
- (c) S(O)NHCH₃, and
- (d) S(O)NHNH₂, and

20

R³ is selected from the group consisting of

- (a) hydrogen,
- (b) CF₃,
- (c) C₁₋₃alkyl and hydroxyC₁₋₃alkyl,
- (d) CN, and

25

the heteroaryl is selected from the group consisting of

30

- (1) 3-isothiazolyl,
- (2) 4-isothiazolyl,
- (3) 5-isothiazolyl,
- (4) 2-oxazolyl,

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5 (5) 4-oxazolyl,
(6) 5-oxazolyl,
(7) 2-thiazolyl,
(8) 4-thiazolyl,
(9) 5-thiazolyl,
(10) 1,2-diazinyl,
(11) 1,3-diazinyl, and
(12) 1,4-diazinyl, and

wherein the substituents are selected from the group consisting of

10 (1) hydrogen,
(2) fluoro or chloro,
(3) methoxy,
(4) methylthio,
(5) CF₃,
(6) methyl.

15

9. A compound according to Claim 1 wherein

X-Y-Z-is selected from the group consisting of:

20 (a) =CH-O-CH=, and
(b) =CH-NR³-CH=,
(c) =N-S-CH=,
(d) =CH-S-N=,
(e) =N-O-CH=,
(f) =CH-O-N=,
25 (g) =N-S-N=,
(h) =N-O-N=,

R¹ is selected from the group consisting of

30 (a) S(O)₂CH₃,
(b) S(O)₂NH₂,
(c) S(O)₂NHC(O)CF₃,
(d) S(O)(NH)CH₃,

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- (e) $\text{S}(\text{O})(\text{NH})\text{NH}_2$, and
- (f) $\text{S}(\text{O})(\text{NH})\text{NHC}(\text{O})\text{CF}_3$;

R^2 is selected from the group consisting of

- (a) $\text{C}_1\text{-4alkyl}$,
- (b) C_3 , C_4 , C_5 , C_6 , and C_7 , cycloalkyl,
- 5 (c) mono- or di-substituted phenyl wherein the substituent is selected from the group consisting of
 - (1) hydrogen,
 - (2) fluoro, chloro, and bromo,
 - (3) $\text{C}_1\text{-4alkoxy}$,
 - (4) $\text{C}_1\text{-4alkylthio}$,
 - (5) CN ,
 - (6) CF_3 ,
 - (7) $\text{C}_1\text{-4alkyl}$,
 - (8) N_3 ,
 - (9) $-\text{CO}_2\text{H}$,
 - (10) $-\text{CO}_2\text{-C}_1\text{-3alkyl}$,
 - (10) $-\text{C}(\text{R}^5)(\text{R}^6)\text{-OH}$ and
 - (11) $-\text{C}(\text{R}^5)(\text{R}^6)\text{-O-C}_1\text{-3alkyl}$,
- 15 (d) mono- or di-substituted heteroaryl selected from the group consisting of
 - (1) furanyl,
 - (2) diazinyl, triazinyl and tetrazinyl,
 - (3) imidazolyl,
 - (4) isooxazolyl,
 - (5) isothiazolyl,
 - (6) oxadiazolyl,
 - (7) oxazolyl,
 - (8) pyrazolyl,
 - (9) pyrrolyl,
 - (10) thiadiazolyl,
 - 20 (11) thiazolyl,
- 25
- 30

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- (12) thienyl,
- (13) triazolyl, and
- (14) tetrazolyl,

wherein said substituents are selected from the group consisting of

- (a) hydrogen,
- (b) fluoro, chloro, bromo,
- (c) C₁-4alkoxy,
- (d) C₁-4alkylthio,
- (e) CN,
- (5) CF₃,
- (6) C₁-4alkyl,
- (7) N₃,
- (8) -C(R⁵)(R⁶)-OH;
- (9) -C(R⁵)(R⁶)-O-C₁-4alkyl.

15 10. A compound according to Claim 9 wherein

R² is selected from the group consisting of

- (a) cyclohexyl, and
- (b) mono or di substituted phenyl, and

20 wherein the substituents are selected from the group
consisting of

- (1) hydrogen,
- (2) halo,
- (3) C₁-4alkoxy,
- (4) C₁-4alkylthio,
- (5) CN,
- (6) CF₃,
- (7) C₁-4alkyl,
- (8) N₃, and
- (9) -C(R⁵)(R⁶)-OH;

25 30 R³ is selected from the group consisting of

- (a) hydrogen,

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- (b) CF₃,
- (c) C₁-3alkyl and hydroxyC₁-3alkyl,
- (d) CN;

R⁵, R^{5'}, R⁶, are each independently selected from the group consisting of

- 5 (a) hydrogen,
- (b) methyl or ethyl,

or R⁵ and R⁶ together with the carbon to which they are attached form a saturated carbon ring of 4, 5 or 6 atoms.

11. A compound according to claim 10 wherein
10 X-Y-Z-is selected from the group consisting of:

- (a) =CH-O-CH=,
- (b) =N-S-N=,
- (c) =N-O-N=;

15 R¹ is selected from the group consisting of
(a) S(O)₂CH₃, and
(b) S(O)₂NH₂;

20 R² is selected from the group consisting of
mono- or di-substituted phenyl wherein the substituents are
selected from the group consisting of

- 20 (1) hydrogen,
- (2) halo, selected from the group consisting of fluoro,
chloro and bromo,
- (3) C₁-3alkoxy,
- (4) C₁-3alkylthio,
- 25 (5) CF₃,
- (6) C₁-3alkyl;

30 R³ is selected from the group consisting of
(a) hydrogen,
(b) CF₃,

(c) C₁-3alkyl and hydroxyC₁-3alkyl,

R⁵ and R⁶ are each selected from the group consisting of

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(a) hydrogen,
(b) methyl or ethyl,
or R⁵, R^{5'} and R⁶ together with the carbon to which they are attached form a saturated carbon ring of 5, 6 or 7 atoms.

5 12. A compound according to claim 11 wherein
X-Y-Z-is =CH-O-CH=;
R¹ is selected from the group consisting of
 (a) S(O)2CH₃, and
 (b) S(O)2NH₂;

10 R² is selected from the group consisting of
 mono- and di-substituted phenyl wherein the substituents are
 selected from the group consisting of
 (1) hydrogen,
 (2) halo, selected from the group consisting of fluoro,
15 chloro and bromo,
 (3) methoxy or ethoxy,
 (4) methyl or ethyl.

13. A compound according to Claim 9 wherein
20 R² is a mono- or di-substituted heteroaryl wherein heteroaryl is
 selected from the group consisting of
 (1) furanyl,
 (2) diazinyl, triazinyl, tetrazinyl,
 (3) imidazolyl,
25 (4) isoxazolyl,
 (5) isothiazolyl,
 (6) oxadiazolyl,
 (7) oxazolyl,
 (8) pyrazolyl,
30 (9) pyrrolyl,
 (10) thiadiazolyl,

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- (11) thiazolyl,
- (12) thiaryl,
- (13) triazolyl,
- (15) pyridyl, and
- (16) tetrazolyl, and

5 wherein the substituents are selected from the group consisting of

- (a) hydrogen,
- (b) fluoro or chloro,
- (c) C₁-3alkoxy,
- (d) C₁-6alkylthio,
- (e) CN,
- (5) CF₃,
- (6) C₁-3alkyl,
- (7) -C(R⁵)(R⁶)-OH;
- (8) -C(R⁵)(R⁶)-O-C₁-4alkyl.

14. A compound according to Claim 13 wherein R¹ is selected from the group consisting of

- (a) S(O)₂CH₃, and
- (b) S(O)₂NH₂, and

20 the hetereoaryl is selected from the group consisting of

- (1) 3-isothiazolyl,
- (2) 4-isothiazolyl,
- (3) 5-isothiazolyl,
- (4) 2-oxazolyl,
- (5) 4-oxazolyl,
- (6) 5-oxazolyl,
- (7) 2-thiazolyl,
- (8) 4-thiazolyl,
- (9) 5-thiazolyl,
- (10) 1,2-diazinyl,

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- (11) 1,3-diazinyl, and
- (12) 1,4-diazinyl, and

wherein the substituents are selected from the group consisting of

- (1) hydrogen,
- (2) fluoro or chloro,
- (3) methoxy,
- (4) methylthio,
- (5) CF₃,
- (6) methyl.

10 15. A compound according to Claim 1 selected from

- (1) 3-(4-(Aminosulfonyl)phenyl)-2-(4-fluorophenyl)-5-(2-hydroxy-2-propyl)thiophene,
- (2) 3-(4-(Aminosulfonyl)phenyl)-2-(4-fluorophenyl)thiophene,
- (3) 3-(4-(Aminosulfonyl)phenyl)-2-(4-fluorophenyl)-5-(2-propyl)thiophene,
- (4) 3-(4-(Aminosulfonyl)phenyl)-2-cyclohexylthiophene,
- (5) 5-(4-Carboxyphenyl)-4-(4-(methylsulfonyl)phenyl)thiophene-2-carboxylic acid,
- (6) 4-(4-Fluorophenyl)-2-methyl-5-(4-(methylsulfonyl)phenyl)thiazole,
- (7) 2-(4-Fluorophenyl)-3-(4-(methylsulfonyl)phenyl)-2-cyclopenten-1-one
- (8) 4-(4-(Methylsulfonyl)phenyl)-5-(4-fluorophenyl)-isothiazole,
- (9) 3-(4-Fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5H)-furanone,
- (10) 3-(4-Fluorophenyl)-4-(4-(aminosulfonyl)phenyl)-2-(5H)-furanone,
- (11) 3-(4-Fluorophenyl)-4-(4-(methylsulfonyl)phenyl)furan,

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(12) 5,5-Dimethyl-3-(4-fluorophenyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(13) 2-(4-(Aminosulfonyl)phenyl)-3-(4-fluorophenyl)thiophene, and
(14) 3-(4-(Trifluoroacetylaminosulfonyl)phenyl)-2-(4-fluorophenyl)thiophene,
5
(15) 3-(2,4-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(16) 3-(3,4-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
10
(17) 3-(2,6-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(18) 3-(2,5-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
15
(19) 3-(3,5-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(20) 3-(4-Bromophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
20
(21) 3-(4-Chlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(22) 3-(4-Methoxyphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
25
(23) 3-(Phenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(24) 3-(2-Chlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
30
(25) 3-(2-Bromo-4-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(26) 3-(2-Bromo-4-Chlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(27) 3-(4-Chloro-2-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,

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- (28) 3-(3-Bromo-4-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (29) 3-(3-Chlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (30) 3-(2-Chloro-4-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (31) 3-(2,4-Dichlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (32) 3-(3,4-Dichlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (33) 3-(2,6-Dichlorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (34) 3-(3-Chloro-4-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (35) 3-(4-Trifluoromethylphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (36) 3-(3-Fluoro-4-methoxyphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (37) 3-(3-Chloro-4-methoxyphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (38) 3-(3-Fluoro-4-methoxyphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (39) 3-(2-Fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (40) 3-(4-Methylthiophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (41) 3-(3-Fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (42) 3-(2-Chloro-6-fluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
- (43) 3-(3-Bromo-4-methylphenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,

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(44) 3-(4-Bromo-2-fluorophenyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(45) 3-(3,4-Dibromophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
5 (46) 3-(4-Chloro-3-fluorophenyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(47) 3-(4-Bromo-3-fluorophenyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(48) 3-(4-Bromo-2-chlorophenyl)-4-(4-methylsulfonyl)phenyl)-2-(5*H*)-furanone,
10 (49) 3-(2-Naphthyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(50) 3-(7-Quinolinyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
(51) 3-(3,4-Dichlorophenyl)-4-(4-(aminosulfonyl)phenyl)-2-(2*H*)-furanone,
15 (52) 3-(3,4-Difluorophenyl)-4-(4-(aminosulfonyl)phenyl)-2-(2*H*)-furanone,
(53) 3-(3-Chloro-4-methoxyphenyl)-4-(4-aminosulfonyl)phenyl)-2-(2*H*)-furanone, and
20 (54) 3-(3-Bromo-4-methoxyphenyl)-4-(4-aminosulfonyl)phenyl)-2-(2*H*)-furanone.

16. A compound which is

25 (a) 3-(3,4-Difluorophenyl)-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone, or
(b) 3-phenyl-4-(4-(methylsulfonyl)phenyl)-2-(5*H*)-furanone,
or a pharmaceutically acceptable salt thereof.

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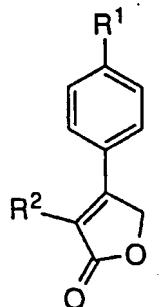
17. A pharmaceutical composition for treating an inflammatory disease susceptible to treatment with a non-steroidal anti-inflammatory agent comprising:
a non-toxic therapeutically effective amount of a compound according
5 to Claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15 or 16.

18. A method of treating an inflammatory disease susceptible to treatment with a non-steroidal anti-inflammatory agent comprising:
administration to a patient in need of such treatment of a non-toxic
10 therapeutically effective amount of a compound according to Claim 1 and a pharmaceutically acceptable carrier.

19. A process of making a compound of formula XXXIII

15

20



XXXIII

25 or a pharmaceutically acceptable salt thereof wherein:
R¹ is selected from the group consisting of
(a) S(O)₂CH₃,
(b) S(O)₂NH₂,
30 (c) S(O)₂NHC(O)CF₃,
(d) S(O)(NH)CH₃,

- 155 -

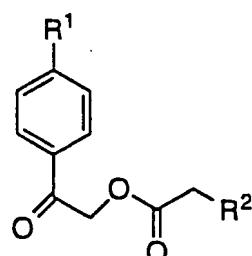
(e) $\text{S}(\text{O})(\text{NH})\text{NH}_2$, and
 (f) $\text{S}(\text{O})(\text{NH})\text{NHC}(\text{O})\text{CF}_3$,

R^2 is selected from the group consisting of
mono- or di-substituted phenyl, and
wherein the substituents are selected from the group

5 consisting of
(1) hydrogen,
(2) halo,
(3) C₁-4alkoxy,
(4) C₁-4alkylthio,
10 (5) CN,
(6) CF₃,
(7) C₁-4alkyl,
(8) N₃, and
(9) -C(R⁵)(R⁶)-O-

15 R^5 and R^6 , are each independently selected from the group consisting of
 (a) hydrogen,
 (b) methyl or ethyl,
 or R^5 and R^6 together with the carbon to which they are attached
 form a saturated carbon ring of 4, 5 or 6 atoms;

20 treating in a non-aqueous polar solvent a compound for formula A



A

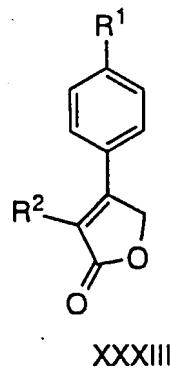
30 in the presence of a strong base:

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to yield a compound of formula XXXIII

5

10



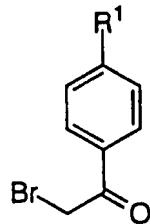
15

20. A process according to claim 19 comprising:

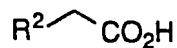
(a) reacting in a non-aqueous polar solvent a compound of formula
XXXII

20

25



with a compound of formula

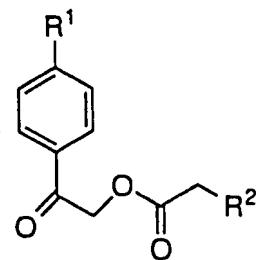


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in the presence of a base to produce a compound of formula A

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5



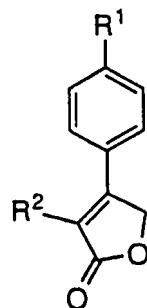
A

10

(b) treating in a non-aqueous polar solvent a compound of formula A
with strong base to yield a compound of formula XXXIII

15

20



XXXIII

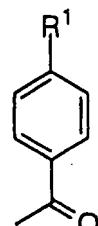
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21. A process according to Claim 20 comprising:
(a1) reacting in an organic solvent a compound of formula XXXII'

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5

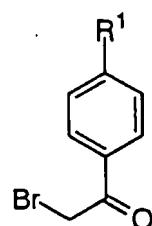


XXXII'

with a bromine reagent to yield a compound of formula XXXII

10

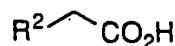
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XXXII

(a2) reacting in a non-aqueous polar solvent a compound of formula
XXXII

20 with a compound of formula



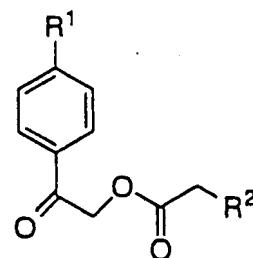
in the presence of a base to produce a compound of formula A

25

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5



A

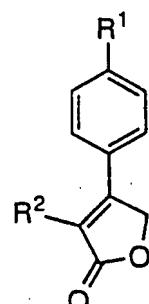
10

(a3) treating in a non-aqueous polar solvent a compound of formula
A

with strong base to yield a compound of formula XXXIII

15

20



XXXIII

25

22. A process according to claim 21 wherein
R¹ is selected from the group consisting of

- (a) S(O)₂CH₃,
- (b) S(O)₂NH₂,
- (c) S(O)NHCH₃, and
- (d) S(O)NHNH₂;

30

R² is

- 160 -

mono or di-substituted phenyl wherein the substitutents are selected from the group consisting of

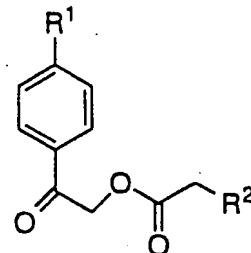
5

- (1) hydrogen,
- (2) halo, selected from the group consisting of fluoro, chloro and bromo,
- (3) methoxy, and
- (4) methyl.

23. A compound of formula A

10

15



A

wherein

R¹ is selected from the group consisting of

20

- (a) S(O)₂CH₃,
- (b) S(O)₂NH₂,
- (c) S(O)₂NHC(O)CF₃,
- (d) S(O)(NH)CH₃,
- (e) S(O)(NH)NH₂, and
- (f) S(O)(NH)NHC(O)CF₃,

25

R² is selected from the group consisting of
mono- or di-substituted phenyl, and
wherein the substitutents are selected from the group
consisting of

30

- (1) hydrogen,
- (2) halo,

- 161 -

- (3) C1-4alkoxy,
- (4) C1-4alkylthio,
- (5) CN,
- (6) CF₃,
- (7) C1-4alkyl,
- 5 (8) N₃, and
- (9) -C(R⁵)(R⁶)-OH;

R⁵ and R⁶, are each independently selected from the group consisting of

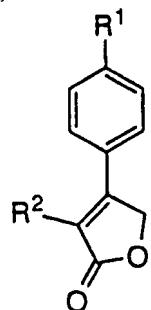
- (a) hydrogen,
- (b) methyl or ethyl,

10 or R⁵ and R⁶ together with the carbon to which they are attached form a saturated carbon ring of 4, 5 or 6 atoms.

24. A process of making a compound of formula XXXIII

15

20



XXXIII

25 wherein

R¹ is selected from the group consisting of

- (a) S(O)₂CH₃,
- (b) S(O)₂NH₂,
- (c) S(O)₂NHC(O)CF₃,
- (d) S(O)(NH)CH₃,
- (e) S(O)(NH)NH₂, and

30

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(f) $\text{S(O)(NH)NHC(O)CF}_3$,

R^2 is selected from the group consisting of
mono- or di-substituted phenyl, and
wherein the substituents are selected from the group

consisting of

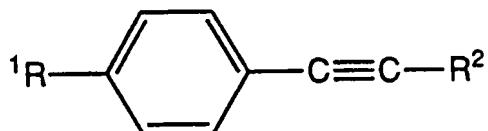
5 (1) hydrogen,
(2) halo,
(3) $\text{C}_1\text{-4alkoxy}$,
(4) $\text{C}_1\text{-4alkylthio}$,
(5) CN,
10 (6) CF_3 ,
(7) $\text{C}_1\text{-4alkyl}$,
(8) N_3 , and
(9) $-\text{C}(\text{R}^5)(\text{R}^6)\text{-OH}$;

R^5 and R^6 , are each independently selected from the group consisting of

15 (a) hydrogen,
(b) methyl or ethyl,
or R^5 and R^6 together with the carbon to which they are attached
form a saturated carbon ring of 4, 5 or 6 atoms.

comprising:

20 (b1) reacting an acetylene compound of the formula XLVIII



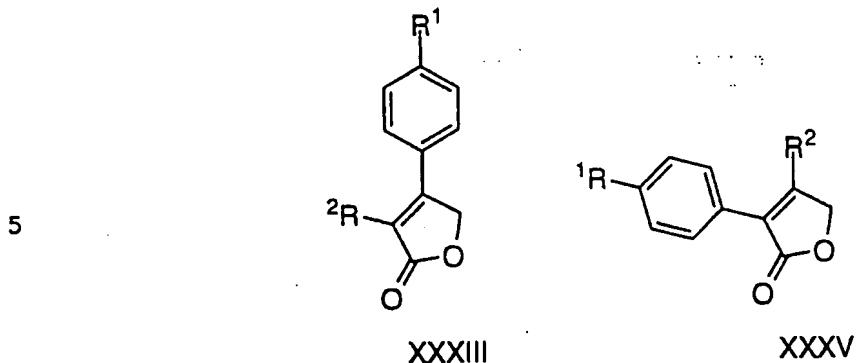
25

XLVIII

with carbon monoxide and water in the presence of a suitable catalyst to
yield a compound of formula XXXIII and XXXV

30

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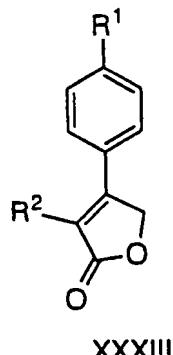


10

25. A process of making a compound of formula XXXIII

15

20



wherein

R¹ is S(O)₂CH₃,R² is selected from the group consisting of

25 mono- or di-substituted phenyl, and

wherein the substituents are selected from the group

consisting of

(1) hydrogen,

(2) halo,

(3) C₁₋₄alkoxy,(4) C₁₋₄alkylthio,

30

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- (5) CN,
- (6) CF₃,
- (7) C₁₋₄alkyl,
- (8) N₃, and
- (9) -C(R⁵)(R⁶)-OH;

5 R⁵ and R⁶, are each independently selected from the group consisting of

- (a) hydrogen,
- (b) methyl or ethyl,

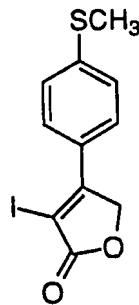
or R⁵ and R⁶ together with the carbon to which they are attached form a saturated carbon ring of 4, 5 or 6 atoms.

10 comprising:

(c1) reacting a compound of formula LIII

15

20



LIII

25 with a reagent of the formula (HO)₂BR² in an aqueous solvent and in the presence of a suitable catalyst to yield a compound of formula LV, and

30

**NOT TO BE TAKEN
INTO CONSIDERATION
FOR THE PURPOSES
OF INTERNATIONAL PROCESSING**

27. A pharmaceutically acceptable salt of a compound of formula (I) as defined in claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15.

5 28. A compound of formula (I) as defined in claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15, or a pharmaceutically acceptable salt thereof for use in treatment of an inflammatory disease susceptible to treatment with a non-steroidal anti-inflammatory agent.

10 29. The compound or salt of claim 16 for use in treatment of an inflammatory disease susceptible to treatment with a non-steroidal anti-inflammatory agent.

15 30. Use of a compound of formula (I) as defined in claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15, or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment of an inflammatory disease susceptible to treatment with a non-steroidal anti-inflammatory agent.

20 31. Use of the compound or salt of claim 16 in the manufacture of a medicament for the treatment of an inflammatory disease susceptible to treatment with a non-steroidal anti-inflammatory agent.

25 32. A non-steroidal anti-inflammatory pharmaceutical composition comprising an acceptable anti-inflammatory amount of a compound of formula (I) as defined in claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15, or a pharmaceutically acceptable salt thereof, in association with a pharmaceutically acceptable carrier.

33. A non-steroidal anti-inflammatory pharmaceutical composition comprising an acceptable anti-inflammatory amount of the compound or salt of claim 16, in association with a pharmaceutically acceptable carrier.